

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
6 May 2005 (06.05.2005)

PCT

(10) International Publication Number
WO 2005/039587 A1

(51) International Patent Classification⁷: **A61K 31/519**,
31/551, C07D 475/08, A61P 37/00, 37/06, 9/00, 25/00,
37/08, 29/00, 31/12, 35/00

(21) International Application Number:
PCT/EP2004/011836

(22) International Filing Date: 18 October 2004 (18.10.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
0324324.3 17 October 2003 (17.10.2003) GB
0408955.3 22 April 2004 (22.04.2004) GB

(71) Applicant (for all designated States except US): **AZA BIOSCIENCE NV** [BE/BE]; Kapucijnenvoer 33, B-3000 Leuven (BE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **WAER, Mark, Jozef, Albert** [BE/BE]; Waversebaan 263, B-3001 Heverlee (BE). **HERDEWIJN, Piet, André, Maurits, Maria**

[BE/BE]; Olivierstraat 21, B-3111 Rotselaar/Wezemaal (BE). **DE JONGHE, Steven, Cesar, Alfons** [BE/BE]; Opaallaan 83, B-1030 Brussel (BE). **MARCHAND, Arnaud, Didier, Marie** [FR/BE]; Armand Thiérylaan 10, B-3001 Heverlee (BE). **GAO, Ling-Jie** [CN/BE]; Leeuwerikenstraat 57/304, B-3001 Heverlee (BE).

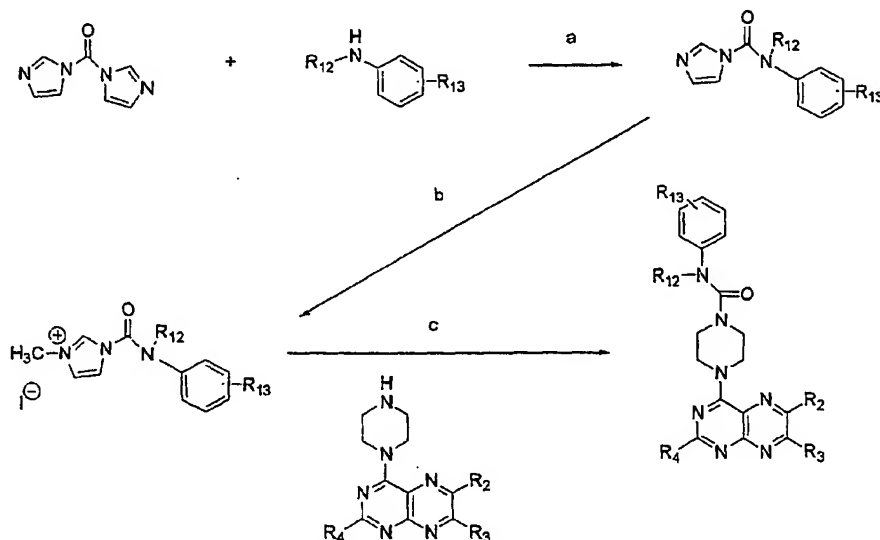
(74) Agents: **BIRD, Arlane et al.**; Bird Goën & Co., Klein Dalenstraat 42A, B-3020 Winksele (BE).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),

[Continued on next page]

(54) Title: HETEROCYCLE-SUBSTITUTED PTERIDINE DERIVATIVES AND THEIR USE IN THERAPY



(57) Abstract: This invention relates to a group of substituted pteridine derivatives, their pharmaceutically acceptable salts, N-oxides, solvates, dihydro- and tetrahydro-derivatives and enantiomers, possessing unexpectedly desirable pharmaceutical properties, in particular which are highly active immunosuppressive agents, and as such are useful in the treatment in transplant rejection and/or in the treatment of certain inflammatory diseases. These derivatives are also useful in preventing or treating cardiovascular disorders, allergic conditions, disorders of the central nervous system, TNF- α related disorders, viral diseases and cell proliferative disorders.



European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

Declaration under Rule 4.17:

— of inventorship (Rule 4.17(iv)) for US only

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

HETEROCYCLE-SUBSTITUTED PTERIDINE DERIVATIVES AND THEIR USE IN THERAPY

FIELD OF THE INVENTION

5 The invention relates to a class of novel pteridines. The invention further relates to pharmaceutical compositions including a broad class of pteridines especially for the prevention and/or the treatment of pathologic conditions such as, but not limited to, immune and auto-immune disorders, organ and cells transplant rejections, cell proliferative disorders, cardiovascular disorders, allergic conditions, disorders of the central nervous system and viral diseases.

10 The invention further relates to combined pharmaceutical preparations comprising one or more such pteridines and one or more known immunosuppressant drugs or antineoplastic drugs or anti-viral drugs.

This invention also relates to a method for the prevention and/or treatment of pathologic conditions such as, but not limited to, immune and autoimmune disorders, organ
15 and cells transplant rejections, cell proliferative disorders, cardiovascular disorders, disorders of the central nervous system, allergic conditions and viral diseases by the administration of an effective amount of a pteridine derivative optionally combined with one or more known immunosuppressant drugs or antineoplastic drugs or anti-viral drugs.

The present invention also relates to the treatment of side effects of various
20 chemotherapeutic drugs and/or of irradiation in cancer therapy. The present invention also relates to the treatment of septic shock, as well as toxic side effects, disorders and diseases related to or resulting from the exposure of patients to abnormally high levels of tumor necrosis factor-alpha (hereinafter referred as TNF- α) in general, and particularly following the administration of TNF- α in cancer treatment in humans. This invention also relates to the
25 treatment of radiotherapy-induced or chemotherapy-induced disorders such as mucositis, secondary myelodysplastic syndromes and radiation-induced graft-versus-host disease, and for the prevention and/or the treatment of injuries in cancer patients such as, but not limited to, apoptosis, radiation necrosis and nephrotoxicity following the administration of certain chemotherapeutic drugs such as cisplatin in cancer treatment. Additionally the invention
30 relates to the treatment of cachexia.

BACKGROUND OF THE INVENTION

Several 2,4-diaminopteridine derivatives being substituted in the 6-position and/or the 7-position of the pteridine ring (according to standard atom numbering for said ring) are known in the art, e.g. from various sources of literature including Swiss Patent No. 231,852; British Patent
35 No. 763,044; United States Patents No. 2,512,572; No. 2,581,889; No. 2,665,275; No. 2,667,486; No. 2,940,972; No. 3,081,230 and No. 5,047,405. Some of these substituted 2,4-diaminopteridine derivatives were disclosed in relationship with various medical uses, such as bacterial growth inhibitors, antineoplastic agents, anti-schistosomiasis activity, coronary dilating

activity, diuretic and hypotensive activity, and anti-amnesic activity. In particular, U.S. Patent No. 2,940,972 and EP-A-362,645 disclose very specific 2,4-diaminopteridine derivatives being substituted by piperidinyl, morpholinyl or pyrrolidinyl in the 7-position of the pteridine ring.

EP-A-185,259 discloses tri- and tetrasubstituted pteridines wherein the substituent in
5 position 2 of the pteridine ring is N-formylpiperazino; the substituent in position 4 of the pteridine ring is selected from the group consisting of dialkylamino, phenylalkylamino, N-alkyl-phenylalkylamino, pyrrolidino, piperidino, (thio)morpholino, 1-oxidothiomorpholino and 1-oxidothioazolidino; the substituent in position 6 of the pteridine ring is selected from the group consisting of hydrogen, alkyl and phenyl; and the substituent in position 7 of the pteridine ring is
10 selected from the group consisting of (di)alkylamino, phenylalkylamino, N-alkyl-phenylalkylamino, piperidino, (thio)morpholino and 1-oxidothiomorpholino. These compounds are suggested for the prophylaxis of thromboembolic disease and arteriosclerosis and for the treatment of tumor growth.

EP-A-574,906 discloses 2,7-diaminopteridines having a *tert*-butoxycarbonylpiperazinyl group in position 4 or in position 6 of the pteridine ring, such compounds being useful for lipid
15 peroxidation inhibition.

Merz et al. in *J. Medicinal Chem.* (1998) 41:4733-4743 discloses 2-N-acetylpiperazino-4-pyrrolidino-6-chloro-7-benzylaminopteridine being useful for inhibiting cAMP phosphodiesterase and malignant tumor cell growth.

WO 02/32507 discloses a series of 7-aminopteridines wherein the substituent in position
20 2 of the pteridine ring may be, among others, SR wherein R is alkyl, cycloalkyl, alkenyl or alkynyl; the substituent in position 4 of the pteridine ring may be, among others, NR^2R^3 wherein R^2 and R^3 are each independently selected from the group consisting of hydrogen, alkyl, cycloalkyl, alkenyl and alkynyl, the latter four groups being optionally substituted; and the substituent in position 6 of the pteridine ring is selected from the group consisting of hydrogen, alkyl and phenyl. These compounds are suggested as modulators of chemokine receptors being useful
25 e.g. for the treatment of asthma, rhinitis, rheumatoid arthritis and the like.

Nevertheless, there still is a need in the art for specific and highly therapeutically active compounds, such as, but not limited to, drugs for treating immune and autoimmune disorders, organ and cells transplant rejections, cell proliferative disorders, cardiovascular
30 disorders, disorders of the central nervous system, allergic conditions and viral diseases. In particular, there is a need in the art to provide immunosuppressive compounds, antineoplastic drugs and anti-viral drugs which are active in a minor dose in order to replace existing drugs having significant side effects and to decrease treatment costs.

Currently used immunosuppressive drugs include antiproliferative agents, such as
35 methotrexate (a 2,4-diaminopteridine derivative disclosed by U.S. Patent No. 2,512,572), azathioprine, and cyclophosphamide. Since these drugs affect mitosis and cell division, they have severe toxic effects on normal cells with high turn-over rate such as bone marrow cells and the gastrointestinal tract lining. Accordingly, marrow depression and liver damage are common side effects of these antiproliferative drugs.

Anti-inflammatory compounds used to induce immunosuppression include adrenocortical steroids such as dexamethasone and prednisolone. The common side effects observed with the use of these compounds are frequent infections, abnormal metabolism, hypertension, and diabetes.

5 Other immunosuppressive compounds currently used to inhibit lymphocyte activation and subsequent proliferation include cyclosporine, tacrolimus and rapamycin. Cyclosporine and its relatives are among the most commonly used immunosuppressant drugs. Cyclosporine is typically used for preventing or treating organ rejection in kidney, liver, heart, pancreas, bone marrow, and heart-lung transplants, as well as for the treatment of
10 autoimmune and inflammatory diseases such as Crohn's disease, aplastic anemia, multiple-sclerosis, myasthenia gravis, uveitis, biliary cirrhosis, etc. However, cyclosporines suffer from a small therapeutic dose window and severe toxic effects including nephrotoxicity, hepatotoxicity, hypertension, hirsutism, cancer, and neurotoxicity.

Additionally, monoclonal antibodies with immunosuppressant properties, such as
15 OKT3, have been used to prevent and/or treat graft rejection. Introduction of such monoclonal antibodies into a patient, as with many biological materials, induces several side-effects, such as dyspnea. Within the context of many life-threatening diseases, organ transplantation is considered a standard treatment and, in many cases, the only alternative to death. The immune response to foreign cell surface antigens on the graft, encoded by the major histo-
20 compatibility complex (hereinafter referred as MHC) and present on all cells, generally precludes successful transplantation of tissues and organs unless the transplant tissues come from a compatible donor and the normal immune response is suppressed. Other than identical twins, the best compatibility and thus, long term rates of engraftment, are achieved using MHC identical sibling donors or MHC identical unrelated cadaver donors. However, such ideal
25 matches are difficult to achieve. Further, with the increasing need of donor organs an increasing shortage of transplanted organs currently exists. Accordingly, xenotransplantation has emerged as an area of intensive study, but faces many hurdles with regard to rejection within the recipient organism.

The host response to an organ allograft involves a complex series of cellular
30 interactions among T and B lymphocytes as well as macrophages or dendritic cells that recognize and are activated by foreign antigen. Co-stimulatory factors, primarily cytokines, and specific cell-cell interactions, provided by activated accessory cells such as macrophages or dendritic cells are essential for T-cell proliferation. These macrophages and dendritic cells either directly adhere to T-cells through specific adhesion proteins or secrete cytokines that
35 stimulate T-cells, such as IL-12 and IL-15. Accessory cell-derived co-stimulatory signals stimulate activation of interleukin-2 (IL-2) gene transcription and expression of high affinity IL-2 receptors in T-cells. IL-2 is secreted by T lymphocytes upon antigen stimulation and is required for normal immune responsiveness. IL-2 stimulates lymphoid cells to proliferate and differentiate by binding to IL-2 specific cell surface receptors (IL-2R). IL-2 also initiates helper

T-cell activation of cytotoxic T-cells and stimulates secretion of interferon- γ which in turn activates cytodestructive properties of macrophages. Furthermore, IFN- γ and IL-4 are also important activators of MHC class II expression in the transplanted organ, thereby further expanding the rejection cascade by enhancing the immunogenicity of the grafted organ. The current model of a T-cell mediated response suggests that T-cells are primed in the T-cell zone of secondary lymphoid organs, primarily by dendritic cells. The initial interaction requires cell to cell contact between antigen-loaded MHC molecules on antigen-presenting cells (hereinafter referred as APC) and the T-cell receptor/CD3 complex on T-cells. Engagement of the TCR/CD3 complex induces CD154 expression predominantly on CD4 T-cells that in turn activate the APC through CD40 engagement, leading to improved antigen presentation. This is caused partly by upregulation of CD80 and CD86 expression on the APC, both of which are ligands for the important CD28 co-stimulatory molecule on T-cells. However, engagement of CD40 also leads to prolonged surface expression of MHC-antigen complexes, expression of ligands for 4-1BB and OX-40 (potent co-stimulatory molecules expressed on activated T-cells). Furthermore, CD40 engagement leads to secretion of various cytokines (e.g., IL-12, IL-15, TNF- α , IL-1, IL-6, and IL-8) and chemokines, all of which have important effects on both APC and T-cell activation and maturation. Similar mechanisms are involved in the development of auto-immune disease, such as type I diabetes. In humans and non-obese diabetic mice, insulin-dependent diabetes mellitus results from a spontaneous T-cell dependent auto-immune destruction of insulin-producing pancreatic β cells that intensifies with age. The process is preceded by infiltration of the islets with mononuclear cells (insulitis), primarily composed of T lymphocytes. A delicate balance between auto-aggressive T-cells and suppressor-type immune phenomena determines whether expression of auto-immunity is limited to insulitis or not. Therapeutic strategies that target T-cells have been successful in preventing further progress of the auto-immune disease. These include neonatal thymectomy, administration of cyclosporine, and infusion of anti-pan T-cell, anti-CD4, or anti-CD25 (IL-2R) monoclonal antibodies. The aim of all rejection prevention and auto-immunity reversal strategies is to suppress the patient's immune reactivity to the antigenic tissue or agent, with a minimum of morbidity and mortality. Accordingly, a number of drugs are currently being used or investigated for their immunosuppressive properties. As discussed above, the most commonly used immunosuppressant is cyclosporine, which however has numerous side effects. Accordingly, in view of the relatively few choices for agents effective at immunosuppression with low toxicity profiles and manageable side effects, there exists a need in the art for identification of alternative immunosuppressive agents and for agents acting as complement to calcineurin inhibition.

The metastasis of cancer cells represents the primary source of clinical morbidity and mortality in the large majority of solid tumors. Metastasis of cancer cells may result from the entry of tumor cells into either lymphatic or blood vessels. Invasion of lymphatic vessels results in metastasis to regional draining lymph nodes. From the lymph nodes, melanoma

cells for example tend to metastasize to the lung, liver, and brain. For several solid tumors, including melanoma, the absence or the presence of lymph nodes metastasis is the best predictor of patient survival. Presently, to our knowledge, no treatment is capable of preventing or significantly reducing metastasis. Hence, there is a need in the art for compounds having such anti-metastasis effect for a suitable treatment of cancer patients.

In the field of allergy, IgE is well known for inducing allergy mainly by stimulating mast cells to release histamine. Also, asthma, being characterized by inflammation of airway and bronchospasm, is mainly induced by Th2 cytokines such as IL-5, IL-10 or IL-13. Therefore there is a need in the art for compounds that efficiently inhibit the release of these Th2 cytokines.

There is also a need in the art to improve therapeutic efficiency by providing pharmaceutical compositions or combined preparations exhibiting a synergistic effect as a result of combining two or more immunosuppressant drugs, or antineoplastic drugs or anti-viral drugs or anti-histamine drugs.

Septic shock is a major cause of death in intensive care units (about 150,000 estimated deaths annually in the United States of America, despite treatment with intravenous antibiotics and supportive care) for which very little effective treatment is available at present. Patients with severe sepsis often experience failures of various systems in the body, including the circulatory system, as well as kidney failure, bleeding and clotting. Lipopolysaccharide (hereinafter referred as LPS) is the primary mediator of Gram-negative sepsis, the most common form of sepsis, by inducing the production of a whole array of macrophage-derived cytokines (such as TNF- α ; interleukins such as IL-1, IL-6, IL-12; interferon-gamma (hereinafter referred IFN- γ), etc.). These cytokines may induce other cells (e.g. T cells, NK cells) to make cytokines as well (e.g. IFN- γ). In addition, other macrophage products (e.g. nitric oxide, hereinafter referred as NO) may also play a role in the pathogenesis of toxic shock. These substances (e.g. NO) may be induced directly due to microbial interactions or indirectly through the action of proinflammatory cytokines. LPS binds to a serum protein known as LPB and the LPS-LPB complex thus formed is recognized by the CD14 toll-like receptor 4 (hereinafter referred as Tlr 4) complex on mononuclear phagocytes. Tlr4 is a signal transducing unit, the activation of which results in the release of mediators such as TNF- α , IL-1 α , IL-1 β and IL-6. These cytokines are important for the pathogenesis of shock. Their administration produces the clinical symptoms of septic shock and their blockade partially protects against LPS-induced lethal shock.

Current therapeutic strategies for the treatment of septic shock are directed against LPS (e.g. antibodies against LPS or LBP-34-23) or against the cytokines induced by LPS (e.g. TNF antibodies) or against the receptor for LPS (e.g. CD14). Unfortunately the initial clinical data of these approaches are very disappointing and illustrate the redundancy of receptors and mediators involved in the pathogenesis of toxic shock. For instance flagellin seems to be

another toxin that plays a role in Gram-negative Salmonella shock syndrome and that cannot be prevented or treated by therapeutic strategies directed specifically at LPS.

Clinical trials in humans with TNF- α blocking antibodies (such as the IL-1 receptor antagonist or PAF receptor antagonists) have been unsuccessful yet, as have been approaches to down regulate inflammation (e.g. using prednisolone) or to block endotoxins. These products must be administered very early after the onset of the disease, which is in most cases not possible.

The only drug currently approved by health authorities for the treatment of adult patients with the most serious forms of sepsis, including septic shock, is a genetically engineered version of a naturally occurring human protein, Activated Protein C, known as Xigris[®] or drotrecogin-alpha which shows only moderate efficacy. Furthermore, because Activated Protein C interferes with blood clotting, the most serious side effect associated with Xigris[®] is bleeding, including bleeding that causes stroke. Thus Xigris[®] is contra-indicated for patients who have active internal bleeding, or who are more likely to bleed because of certain medical conditions including recent strokes, recent head or spinal surgery or severe head trauma. Because treatment with Xigris[®] comes with potentially serious risks, the benefits and risks of treatment with Xigris[®] must be carefully weighed for each individual patient.

Therefore there is a strong need in the art for new medications, either alone or in combination with the currently suggested treatments, for treating the most serious forms of life-threatening illnesses caused by severe infection, such as septic shock.

TNF- α is generally considered to be the key mediator in the mammalian response to bacterial infection. It is a strong pro-inflammatory agent that will affect the function of almost any organ system, either directly or by inducing the formation of other cytokines like IL-1 or prostaglandines. TNF- α is also a potent anti-tumor agent. If administered in small quantities to humans, it causes fever, headache, anorexia, myalgia, hypotension, capillary leak syndrome, increased rates of lipolysis and skeletal muscle protein degradation (including cachexia). Its use in cancer treatment is therefore very much limited by its severe side effects.

TNF- α , a pleiotropic cytokine produced mainly by activated macro-phages, exerts an *in vitro* cytotoxic action against transformed cells and *in vivo* anti-tumor activities in animal models. However, despite the fact that TNF- α is used in cancer patients especially to treat melanoma and sarcoma, the major problem hampering its use is toxicity. Indeed, TNF- α induces shock-like symptoms such as bowel swelling and damage, liver cell necrosis, enhanced release of inflammatory cytokines such as IL-1 or IL-6, and hypo-tension probably due to the release of inducers of vessels dilatation such nitric oxide and other proinflammatory cytokines. Cardiovascular toxicity is usually dose-limiting. Hypotension can be severe with systolic blood pressure below 60 mm Hg. Respiratory compromise is common after treatment with TNF- α and may require mechanical ventilation. Upper as well as lower digestive tract symptoms are also common in this type of treatment. Nausea and vomiting can be distressing

and in some cases dose-limiting. Watery diarrhea is frequently observed. Neurological sequelae of treatment with TNF- α can also occur.

Hence, compounds that inhibit the toxic effects of TNF- α but that do not inhibit TNF- α anti-tumor effect are highly desirable for the treatment of cancer patients. Presently, several clinical trials involving TNF- α are being developed for the cancer of organs such as liver, lung, kidney and pancreas, which are based on a procedure including the steps of organ isolation, injection of TNF- α into the isolated organ, and reperfusion of the treated organ. However, even for isolated organ perfusion, some TNF- α usually escapes to the general blood circulation and leads to the mortality of about 10% of the patients thus treated. Many patients treated by this procedure also require intensive care unit rescue to cope with the toxic side-effects of such TNF- α treatment.

Combined treatment of TNF- α with alkylating drugs in an isolated organ perfusion model has received considerable attention. TNF- α is currently successfully used in isolated limb perfusion of human cancer patients and, in combination with melphalan and interferon-gamma, against melanoma, sarcomas and carcinomas.

The gastrointestinal mucosa is very sensitive to chemotherapeutic drugs. Mucositis caused by chemotherapy usually begins rapidly after initiation of the treatment with inflammation and ulceration of the gastrointestinal tract and leading to diarrhea. Severe, potentially life-threatening, diarrhea may require interruption of the chemotherapeutic treatment and subsequent dose reduction of the therapeutic agent. The oral cavity is often the place of severe side effects from cancer therapy that adversely affects the quality of life of the patient and its ability to tolerate the therapy. These side effects can be caused by radiotherapy as well as chemotherapy. A relationship between both serum and mucosal levels of TNF- α and IL-1 correlates with nonhematologic toxicities, including mucositis.

Radiation injuries occurring e.g. after a single high-dose irradiation include apoptosis as well as radiation necrosis. Even normal tissues protected by shielding during irradiation may be considerably damaged. It was found in experimental animal models that the radiation injuries after a single high-dose irradiation typically used for the treatment of various malignant tumors consist of radiation necrosis and apoptosis, which were correlated with the expression of TNF- α and TGF- β 1.

Irradiation may induce graft-versus-host disease (hereinafter referred as GVHD) in cancer patients. This disease may occur especially in patients receiving allogeneic bone marrow transplantation as a treatment for cancers such as leukemia or lymphoma and can lead to the death of about 25% of the relevant patients. Before bone marrow transplantation, leukaemia patients for example receive either total body or total lymphoid irradiation to suppress their immune system. However, such irradiation induces not only necrosis but also the release of proinflammatory cytokines mainly TNF- α , IL-1 and IL-6 which in turn induce

direct host tissues inflammation and activation of donor cells against host antigens leading to GVHD.

Cisplatin is an effective chemotherapeutic agent used in the treatment of a wide variety of both pediatric and adult malignancies, including testicular, germ cell, head and neck (cervical), bladder and lung cancer. Dose-dependent and cumulative nephrotoxicity is the major side effect of cisplatin, sometimes requiring a reduction in dose or discontinuation of the treatment. Other side effects of cisplatin include kidney damage, loss of fertility, harmful effect on a developing baby, temporary drop in bone marrow function causing drop in white blood cell count, anaemia, drop in platelets causing bleeding, loss of appetite, numbness or tingling in limbs, loss of taste, allergic reactions, and hearing disorders (difficulty in hearing some high-pitched sounds, experiencing ringing in the ears). Blurred vision may also be a side effect with high doses of cisplatin. It was shown that TNF- α is a key element in a network of proinflammatory chemokines and cytokines activated in the kidney by cisplatin. Blockade of TNF- α action would prevent the activation of this cytokine network and would provide protection against cisplatin nephrotoxicity. Hence, compounds that inhibit the toxic effects of cisplatin but that do not inhibit cisplatin anti-tumor effects are highly desirable for the treatment of cancer patients.

A surplus of TNF- α also causes a dramatic change of endothelial cells. In particular, TNF- α is an important mediator of skeletal muscle degeneration associated with cachexia, a debilitating syndrome characterized by extreme weight loss and whole-body wasting. Cachexia is usually a secondary condition whereby there is excessive tissue catabolism in combination with deficient anabolism. It is frequently seen in patients afflicted with chronic diseases such as cancer, cardiopulmonary diseases, aging, malabsorptive disorders, excessive physical stress, eating disorders and acquired immuno-deficiency syndrome (AIDS). Some authors consider that the elevated TNF- α values found in at least 50% of cancer patients in the active stage of the disease can result in cachexia. TNF- α levels in clinically healthy adults, as well as in adult cancer patients, are well documented, for instance by Nenova et al. in *Archives of Hellenic Medicine* (2000) 17:619-621. Serum TNF- α concentrations in healthy children as well as in children with malignancies are documented for instance by Saarinen et al. in *Cancer Research* (1990) 50:592-595. A very significant proportion of cancer mortalities result from cachexia rather than from tumor burden. Chronic wasting disease (cachexia) may result when excessive cellular damage results in the release of substances (TNF- α , collagenase, hyaluronidase) that further catabolize the so-called healthy tissue resulting in an inability to assimilate nutrients required for anabolic restructuring of associated tissue.

Infants infected with human immunodeficiency virus type 1 (HIV-1) show growth retardation and severe weight loss that can lead to death. The overproduction of certain cytokines has been implicated as a possible cause for this. For instance, according to Rautonen et al. in *AIDS* (1991) 5:1319-1325, serum IL-6 concentrations are elevated and

associated with elevated TNF- α concentrations in children with HIV infection. Swapan et al. in *Journal of Virology* (2002) 76:11710-11714 have shown that reduction of TNF- α levels by either anti-TNF- α antibodies or human chorionic gonadotropin inhibits the expression of HIV-1 proteins and prevents cachexia and death.

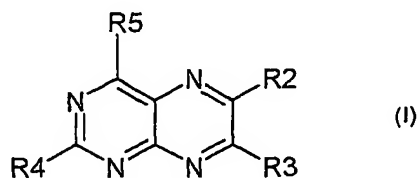
- 5 Very few drugs have been suggest at present for the treatment of cachexia. Some high-dose progestins like megestrol acetate, an agent used for the treatment of metastatic breast cancer, and medroxyprogesterone acetate were shown in randomized clinical trials to provide a statistically significant advantage as regards improved appetite and body weight gain. Hence, compounds that stimulate appetite and body weight gain without inhibiting the
10 anti-tumor effect or anti-viral effect of co-administered drugs are highly desirable for the treatment of cachexia. More specifically, there is a need in the art for treating cachexia by the administration of compounds that reduce TNF- α levels in the serum of humans.

- TNF- α is also suspected to play a role, through a possible dual action in the hematopoietic environment, in the development of hematologic malignancies such as
15 idiopathic myelodysplastic syndromes occurring most often in elderly people but also occasionally in children, these syndromes being currently regarded as the early phase of acute leukemia.

- There is a strong need in the art to improve, or to provide alternatives to, the existing prophylactic or therapeutic solutions to all the aforesaid diseases. Meeting these various
20 needs in the art constitutes the main goal of the present invention.

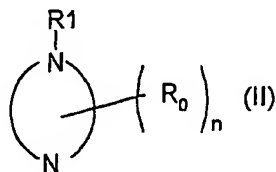
SUMMARY OF THE INVENTION

In a first embodiment, the present invention relates to a group of novel pteridine derivatives having the general formula (I):

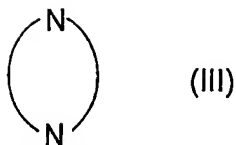


wherein:

- 25 - a first group of one or more of the substituents R_2 , R_3 , R_4 and R_5 of the pteridine ring is independently selected from groups represented by the general formula (II):



wherein:



schematically represents a saturated or partly unsaturated heterocyclic ring with at least two nitrogen atoms in the said heterocyclic ring and with a total of 5 to 7 atoms in the said heterocyclic ring, and optionally with one or more other heteroatoms (e.g. oxygen or sulfur) in the said heterocyclic ring or attached to one or more carbon atoms of said heterocyclic ring (for instance in the form of a carbonyl or thiocarbonyl group), wherein one of said at least two nitrogen atoms in the heterocyclic ring is attached to a carbon atom of the pteridine ring at any of positions 2, 4, 6 or 7 of the pteridine ring, wherein the said heterocyclic ring may be fused to one or more aromatic hydrocarbon rings, and wherein:

- each substituent R_0 of the heterocyclic ring (III) is a group independently selected from the group consisting of halogen, nitro, C_{1-7} alkyl (optionally containing one or more functions or radicals selected from the group consisting of halogen, carbonyl, thiocarbonyl, hydroxyl, sulfhydryl, C_{1-7} alkoxy, thio C_{1-7} alkyl, thio C_{3-10} cycloalkyl, acetal, thioacetal, imino, oximino, alkyloximino, amino-acid, cyano, (thio)carboxylic acid, (thio)carboxylic acid ester or amide, nitro, amino, C_{1-7} alkylamino, cycloalkylamino, alkenylamino, cycloalkenyl-amino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercapto-alkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl and sulfonamido), C_{3-7} alkenyl, C_{2-7} alkynyl, halo C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl, arylalkyl, alkylaryl, alkylacyl, arylacyl, hydroxyl, sulfhydryl, amino, C_{1-7} alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkyl-amino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, C_{1-7} alkoxy, C_{3-10} cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C_{1-7} alkyl, thio C_{3-10} cycloalkyl, thioaryl, thioheterocyclic, arylalkylthio, heterocyclic-substituted alkylthio, formyl, hydroxylamino, cyano, (thio)carboxylic acid or esters or thioesters or amides or thioamides thereof;
- n is an integer from 0 to 6;
- R_1 is a substituent group selected from the group consisting of formyl, acyl, thio-acyl, amide, thioamide, sulfonyl, sulfinyl, carboxylate, thiocarboxylate, amino-substituted acyl, alkoxyalkyl, C_{3-10} cycloalkyl-alkyl, C_{3-10} cyclo-alkyl, dialkylaminoalkyl, heterocyclic-substituted alkyl, acyl-substituted alkyl, thioacyl-substituted alkyl, amido-substituted alkyl, thioamido-substituted alkyl, carboxylato-substituted alkyl, thiocarboxylato-substituted alkyl, (amino-substituted acyl)alkyl, heterocyclic, carboxylic acid ester, ω -cyanoalkyl, ω -carboxylic ester-alkyl, halo C_{1-7} alkyl, C_{2-7}

- alkenyl, C₂₋₇ alkynyl, arylalkenyl, aryloxyalkyl, arylalkyl and aryl, wherein the aryl moiety of each of said arylalkenyl, aryloxyalkyl, arylalkyl and aryl radicals is optionally substituted with one or more substituents independently selected from the group consisting of halogen, C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, halo C₁₋₇ alkyl, nitro, hydroxyl, sulfhydryl, amino, C₁₋₇ alkoxy, C₃₋₁₀ cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, thioaryl, thio-heterocyclic, arylalkylthio, heterocyclic-substituted alkylthio, formyl, carbamoyl, thiocarbamoyl, ureido, thioureido, sulfon-amido, hydroxylamino, alkoxyamino, mercaptoamino, thioalkyl-amino, acylamino, thioacylamino, cyano, carboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, thiocarboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, alkylamino, cycloalkylamino, alkenyl-amino, cycloalkenylamino, alkynylamino, arylamino, aryl-alkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic amino, hydrazino, alkylhydrazino and phenylhydrazino; and
- wherein the remaining of, i.e. a second group of, the substituents R₂, R₃, R₄ and R₅ of the pteridine ring is independently selected from the group consisting of hydrogen; halogen; C₁₋₇ alkyl; C₂₋₇ alkenyl; C₂₋₇ alkynyl; halo C₁₋₇ alkyl; carboxy C₁₋₇ alkyl; carboxyaryl; C₁₋₇ alkoxy; C₃₋₁₀ cycloalkoxy; aryloxy; arylalkyloxy; oxyheterocyclic; heterocyclic-substituted alkyloxy; thio C₁₋₇ alkyl; thio C₃₋₁₀ cycloalkyl; thioaryl; thioheterocyclic; arylalkylthio; heterocyclic-substituted alkylthio; hydroxylamino; mercapto-amino; acylamino; thio-acylamino; alkoxyamino; thioalkyl-amino; acetal; thio-acetal; carboxylic acid; carboxylic acid esters, thioesters, halides, anhydrides, amides and thioamides; thiocarboxylic acid; thiocarboxylic acid esters, thioesters, halides, anhydrides, amides and thioamides; hydroxyl; sulfhydryl; nitro; cyano; carbamoyl; thiocarbamoyl; ureido; thioureido; amino; alkylamino; cycloalkylamino; alkenylamino; cycloalkenylamino; alkynylamino; arylamino; arylalkylamino; hydroxyalkylamino; mercaptoalkyl-amino; heterocyclic amino; heterocyclic-substituted arylamino; heterocyclic-substituted alkyl-amino; oximino; alkyloximino; hydrazino; alkylhydrazino; phenylhydrazino; cysteinyl acid, esters, thioesters, halides, anhydrides, amides and thioamides thereof; aryl optionally substituted with one or more substituents selected from the group consisting of halogen, C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, halo C₁₋₇ alkyl, nitro, hydroxyl, sulfhydryl, amino, C₁₋₇ alkoxy, C₃₋₁₀ cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, thioaryl, thioheterocyclic, arylalkylthio, heterocyclic-substituted alkylthio, formyl, carbamoyl, thiocarbamoyl, ureido, thio-ureido, sulfonamido, hydroxylamino, alkoxyamino, mercaptoamino, thioalkylamino, acylamino, thioacylamino, cyano, carboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, thiocarboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkylamino, hydroxyalkyl-amino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino and phenylhydrazino; optionally

substituted heterocyclic radicals; aromatic or heterocyclic substituents substituted with an aliphatic spacer between the pteridine ring and the aromatic or heterocyclic substituent, whereby said aliphatic spacer is a branched or straight, saturated or unsaturated aliphatic chain of 1 to 4 carbon atoms which may contain one or more functions, atoms or radicals

5 independently selected from the group consisting of carbonyl, thiocarbonyl, hydroxyl, thiol, ether, thioether, acetal, thioacetal, amino, imino, oximino, alkyloximino, amino-acid, cyano, acylamino, thioacylamino, carbamoyl, thiocarbamoyl, ureido, thioureido, carboxylic acid or ester or thioester or halide or anhydride or amide, thiocarboxylic acid or ester or thioester or halide or anhydride or amide, nitro, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, hydroxylamino,

10 mercaptoamino, alkylamino, cycloalkylamino, alkenylamino, cycloalkenyl-amino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl, sulfinyl, sulfonamido and halogen; branched or straight, saturated or unsaturated aliphatic chains of 1 to 7 carbon atoms

15 optionally containing one or more functions, atoms or radicals independently selected from the group consisting of halogen, carbonyl, thiocarbonyl, hydroxyl, thiol, ether, thio-ether, acetal, thio-acetal, amino, imino, oximino, alkyloximino, aminoacid, cyano, acylamino; thioacylamino; carbamoyl, thiocarbamoyl, ureido, thio-ureido, carboxylic acid ester or halide or anhydride or amide, thiocarboxylic acid or ester or thioester or halide or anhydride or amide, nitro, thio C₁₋₇

20 alkyl, thio C₃₋₁₀ cycloalkyl, hydroxylamino, mercaptoamino, alkyl-amino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkyl-amino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, hetero-cyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl, sulfinyl and sulfonamido; or R₂ together with R₃ and the carbon atoms in positions 6

25 and 7 of the pteridine ring forms a homocyclic or heterocyclic radical; with the first proviso that when R₄ stands for N-formylpiperazino then R₂ may not be hydrogen, C₁₋₇ alkyl or phenyl; and with the second proviso that when R₄ stands for N-acetylpiperazino then R₂ may not be halogen;

and/or a pharmaceutically acceptable addition salt thereof and/or a stereoisomer thereof

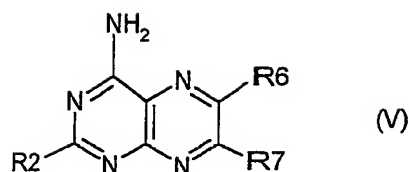
30 and/or a mono- or a di-N-oxide thereof and/or a solvate thereof and/or a dihydro- or tetrahydropteridine derivative thereof.

The above novel compounds of this first embodiment have in common the structural features present in the general formula (I), in particular the pteridine ring is substituted by at least one N,N-containing heterocyclic group being itself N-substituted by a carbonyl or

35 thiocarbonyl or sulfonyl radical or by certain hydrocarbonyl radicals other than C₁₋₇ alkyl. They also have a potential specific biological activity profile and consequent usefulness in medicinal chemistry.

In a second embodiment, the present invention relates to a group of novel 4-amino-pteridine derivatives having the general formula (V):

13



wherein:

- R₂ is selected from the group consisting of nitrogen-containing heterocyclic radicals other than morpholinyl and piperazinyl, said radicals being attached to the pteridine ring by means of a nitrogen atom; benzylamino; phenylethylamino; heterocyclic-substituted alkylamino; aryloxy; arylthio; arylsulfonyl; arylalkyloxy; arylalkylthio; C₁₋₇ alkylsulfonyl; heterocyclic-substituted alkyloxy; and heterocyclic-substituted alkylthio;
- R₆ and R₇ are independently selected from the group consisting of hydrogen; halogen; C₁₋₇ alkyl; C₂₋₇ alkenyl; C₂₋₇ alkynyl; halo C₁₋₇ alkyl; carboxy C₁₋₇ alkyl; C₁₋₇ alkylsulfonyl; carboxyaryl; C₁₋₇ alkoxy; C₃₋₁₀ cycloalkoxy; aryloxy; arylalkyloxy; oxyheterocyclic; heterocyclic-substituted alkyloxy; C₁₋₇ alkylthio; thio C₃₋₁₀ cycloalkyl; arylthio; arylsulfonyl; thio-heterocyclic; arylalkylthio; heterocyclic-substituted alkylthio; hydroxylamino; mercaptoamino; acylamino; thioacylamino; alkoxyamino; thioalkylamino; acetal; thio-acetal; carboxylic acid; carboxylic acid esters, thioesters, halides, anhydrides, amides and thioamides; thiocarboxylic acid; thiocarboxylic acid esters, thioesters, halides, anhydrides, amides and thioamides; hydroxyl; sulfhydryl; nitro; cyano; carbamoyl; thiocarbamoyl; ureido; thioureido; amino; alkyl-amino; cycloalkylamino; alkenylamino; cycloalkenylamino; alkynyl-amino; arylamino; arylalkylamino; hydroxyalkylamino; mercaptoalkyl-amino; heterocyclic amino; heterocyclic-substituted alkylamino; oximino; alkylloximino; hydrazino; alkylhydrazino; phenylhydrazino; cysteinyl acid, esters, thioesters, halides, anhydrides, amides and thioamides thereof; aryl optionally substituted with one or more substituents independently selected from the group consisting of halogen, C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, halo C₁₋₇ alkyl, nitro, hydroxyl, sulfhydryl, amino, C₁₋₇ alkoxy, C₃₋₁₀ cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, thioaryl, thio-heterocyclic, arylalkylthio, heterocyclic-substituted alkylthio, formyl, C₁₋₇ alkanoyl (acyl), carbamoyl, thiocarbamoyl, ureido, thioureido, sulfonamido, hydroxyl-amino, alkoxyamino, mercaptoamino, thioalkylamino, acylamino, thioacylamino, cyano, carboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, thiocarboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic amino, hydrazino, alkylhydrazino and phenylhydrazino; optionally substituted heterocyclic radicals; aryl or heterocyclic radicals substituted with an aliphatic spacer between the pteridine ring and said aryl or heterocyclic radical, whereby said aliphatic spacer is a

5 branched or straight, saturated or unsaturated aliphatic chain of 1 to 4 carbon atoms which may contain one or more functions, atoms or radicals independently selected from the group consisting of carbonyl, thiocarbonyl, hydroxyl, thiol, ether, thioether, acetal, thioacetal, amino, imino, oximino, alkyloximino, amino-acid, cyano, acylamino, thioacyl-amino, carbamoyl, thiocarbamoyl, ureido, thio-ureido, carboxylic acid or ester or thioester or halide or anhydride or amide, thiocarboxylic acid or ester or thioester or halide or anhydride or amide, nitro, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, hydroxylamino, mercaptoamino, alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic amino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl, sulfinyl, sulfonamido and halogen; branched or straight, saturated or unsaturated aliphatic chains of 1 to 7 carbon atoms optionally containing one or more functions, atoms or radicals independently selected from the group consisting of halogen, carbonyl, thiocarbonyl, hydroxyl, thiol, ether, thio-ether, acetal, thio-acetal, amino, imino, oximino, alkyloximino, aminoacid, cyano, acylamino, thioacylamino, carbamoyl, thiocarbamoyl, ureido, thioureido, carboxylic acid ester or halide or anhydride or amide, thiocarboxylic acid or ester or thioester or halide or anhydride or amide, nitro, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, hydroxylamino, mercapto-amino, alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic amino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl, sulfinyl and sulfonamido; or R₆ together with R₇ and the carbon atoms in positions 6 and 7 of the pteridine ring forms a homocyclic or heterocyclic radical;

25 and/or a pharmaceutically acceptable addition salt thereof and/or a stereoisomer thereof and/or a mono- or a di-*N*-oxide thereof and/or a solvate thereof and/or a dihydro- or tetrahydropteridine derivative thereof.

30 The novel compounds of this second embodiment have in common the structural features present in the general formula (V), in particular they are amino-substituted at position 4 of the pteridine ring and substituted at position 2 of the pteridine ring by a radical being a nitrogen-containing or oxygen-containing or sulfur-containing nucleophile. They also have a potential specific biological activity profile and consequent usefulness in medicinal chemistry as will be detailed below. Furthermore, some of the novel compounds of this second embodiment are intermediates for making novel pteridine derivatives having the general formula (I) in the first embodiment, as shown for instance in figures 6 and 7.

35 In a third embodiment, the present invention relates to the unexpected finding that at least one desirable biological property such as, but not limited to, the ability to decrease the proliferation of lymphocytes, or to decrease T-cell activation, or to decrease B-cell or monocytes or macrophages activation, or to inhibit the release of certain cytokines, or in inhibiting human TNF- α production is a feature which is present in the said group of novel

compounds. As a consequence, the invention relates to pharmaceutical compositions comprising as an active principle at least one pteridine derivative having the general formula (I), and/or at least one 4-amino-pteridine derivative having the general formula (V) and/or a pharmaceutically acceptable addition salt thereof and/or a stereoisomer thereof and/or a mono- or a di-N-oxide thereof and/or a solvate and/or a dihydro- or tetrahydropteridine derivative thereof.

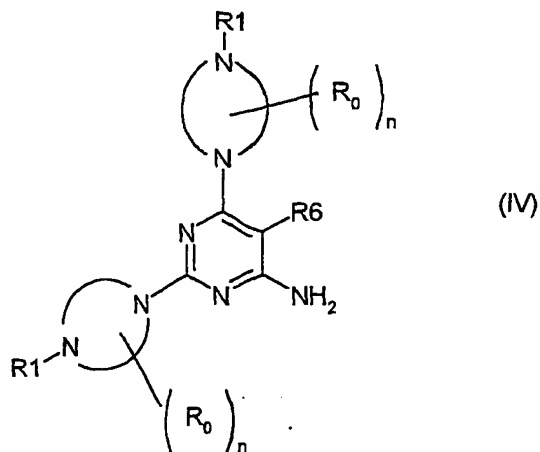
Compounds having the general formulae (I) and (V) are highly active immunosuppressive agents, antineoplastic agents, anti-allergic agents or anti-viral agents which, together with one or more pharmaceutically acceptable carriers, may be formulated into pharmaceutical compositions for the prevention or treatment of pathologic conditions such as, but not limited to, immune and autoimmune disorders, organ and cells transplant rejections, allergic conditions, cell proliferative disorders, cardiovascular disorders, disorders of the central nervous system and viral diseases. Compounds having the general formulae (I) and (V) are also useful for the prevention or treatment of a TNF- α -related disorder in a mammal, such as for instance:

- septic or endotoxic shock,
- TNF- α - mediated diseases,
- pathologies and conditions associated with and/or induced by abnormal levels of TNF- α occurring in a systemic, localized or particular tissue type or location in the body of the mammal,
- toxic effects of TNF- α and/or anti-cancer chemotherapeutic agents,
- injuries after irradiation of a tissue of the mammal by radio-elements, and
- cachexia.

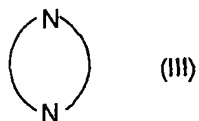
In a further embodiment, the present invention relates to combined preparations containing at least one compound of the general formula (I) or the general formula (V) and one or more drugs such as immunosuppressant and/or immunomodulator drugs, antineoplastic drugs, anti-histamines, inhibitors of agents causative of allergic conditions, or antiviral agents. In a further embodiment, the present invention relates to the prevention or treatment of the above-cited pathologic conditions by administering to the patient in need thereof an effective amount of a compound of the general formula (I) or the general formula (V), optionally in the form of a pharmaceutical composition or combined preparation with another suitable drug.

In another embodiment, the present invention relates to various processes and methods for making the novel pteridine derivatives defined in general formulae (I) and (V), as well as their pharmaceutically acceptable salts, N-oxides, solvates, enantiomers and dihydro- and tetrahydroderivatives.

In a still further embodiment, the present invention relates to a family of novel polysubstituted 6-aminopyrimidines having the general formula (IV):

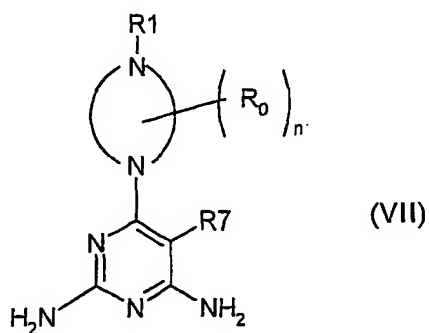


wherein each of n , R_0 and

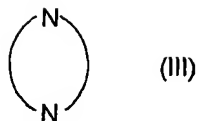


are as defined hereinabove with respect to formula (II); wherein R_1 is as defined hereinabove with respect to formula (II) or is hydrogen; and wherein R_6 is selected from the group consisting of nitro and amino;

In another embodiment, the present invention relates to a family of novel polysubstituted 2,6-diaminopyrimidines having the general formula (VII)



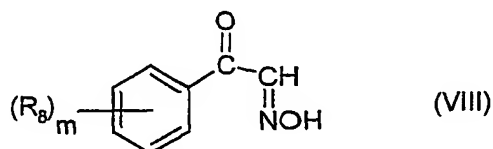
wherein:



and R_0 , R_1 and n are as defined hereinabove with respect to formula (II), and wherein R_7 is selected from the group consisting of hydrogen, nitroso and amino.

The said novel polysubstituted pyrimidines having the general formulae (IV) and (VII) are useful as intermediates for making some of the pteridine derivatives of the present invention.

In another embodiment, the present invention relates to a family of novel substituted phenylglyoxalmonoximes having the general formula (VIII):



wherein m is from 0 to 5, and wherein each substituent R_8 is independently selected from the group consisting of halogen, cyano, piperidino, imidazol-1-yl, hydroxy, amino, protected amino (such as acetamido), nitro, benzoxy, acetoxy, C_{1-7} alkoxy and C_{1-7} alkyl, as well as a method for making them from R_8 -substituted acetophenones. The said novel substituted phenylglyoxal-monoximes having the general formula (VIII) are useful as intermediates for making some of the pteridine derivatives defined in the general formulae (I) and (V) of the present invention.

In yet another embodiment, the present invention relates to a family of novel 2-substituted-4,6-diamino-5-nitrosopyrimidines and 2-substituted-4,5,6-triaminopyrimidines, as well as a method for making them, which are useful as intermediates for making some of the pteridine derivatives of the present invention.

20 BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 schematically shows a first method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II).

Figure 2 schematically shows a second method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II).

Figure 3 schematically shows a third method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₅ is a group having the formula (II).

Figure 4 schematically shows a fourth method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₅ is a group having the formula (II).

Figure 5 schematically shows a fifth method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₅ is a group having the formula (II).

Figure 6 schematically shows a sixth method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₅ is a group having the formula (II).

Figure 7 schematically shows a seventh method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₅ is a group having the formula (II).

Figure 8 schematically shows a method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₄ and R₅ are identical groups having the formula (II).

Figure 9 schematically shows another method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₅ is a specific group having the formula (II).

DEFINITIONS

Unless otherwise stated herein, the term " trisubstituted " means that three of the carbon atoms being in positions 2, 4 and 6 or, alternatively, in positions 2, 4 and 7 of the pteridine ring (according to standard atom numbering for the pteridine ring) are substituted with an atom or group other than hydrogen. The term " tetrasubstituted " means that all four carbon atoms being in positions 2, 4, 6 and 7 of the pteridine ring are substituted with an atom or group other than hydrogen.

As used herein with respect to a substituting radical, and unless otherwise stated, the term " C₁₋₇ alkyl " means straight and branched chain saturated acyclic hydrocarbon monovalent radicals having from 1 to 7 carbon atoms such as, for example, methyl, ethyl, propyl, n-butyl, 1-methylethyl (isopropyl), 2-methylpropyl (isobutyl), 1,1-dimethylethyl (tert-butyl), 2-methylbutyl, n-pentyl, dimethylpropyl, n-hexyl, 2-methylpentyl, 3-methylpentyl, n-heptyl and the like.

As used herein with respect to a substituting radical, and unless otherwise stated, the term " acyl " broadly refers to a carbonyl (oxo) group adjacent to a C₁₋₇ alkyl radical, a C₃₋₁₀ cycloalkyl radical, an aryl radical, an arylalkyl radical or a heterocyclic radical, all of them being such as herein defined; representative examples include acetyl, benzoyl, naphthoyl and

the like; similarly, the term " thioacyl " refers to a C=S (thioxo) group adjacent to one of the said radicals.

As used herein with respect to a substituting radical, and unless otherwise stated, the term " C₁₋₇ alkylene " means the divalent hydrocarbon radical corresponding to the above defined C₁₋₇ alkyl, such as methylene, bis(methylene), tris(methylene), tetramethylene, hexamethylene and the like.

As used herein with respect to a substituting radical, and unless otherwise stated, the term " C₃₋₁₀ cycloalkyl " means a mono- or polycyclic saturated hydrocarbon monovalent radical having from 3 to 10 carbon atoms, such as for instance cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl and the like, or a C₇₋₁₀ polycyclic saturated hydrocarbon monovalent radical having from 7 to 10 carbon atoms such as, for instance, norbornyl, fenchyl, trimethyltricycloheptyl or adamantyl.

As used herein with respect to a substituting radical, and unless otherwise stated, the term " C₃₋₁₀ cycloalkyl-alkyl " refers to an aliphatic saturated hydrocarbon monovalent radical (preferably a C₁₋₇ alkyl such as defined above) to which a C₃₋₁₀ cycloalkyl (such as defined above) is already linked such as, but not limited to, cyclohexylmethyl, cyclopentylmethyl and the like.

As used herein with respect to a substituting radical, and unless otherwise stated, the term " C₃₋₁₀ cycloalkylene " means the divalent hydrocarbon radical corresponding to the above defined C₃₋₁₀ cycloalkyl.

As used herein with respect to a substituting radical, and unless otherwise stated, the term " aryl " designate any mono- or polycyclic aromatic monovalent hydrocarbon radical having from 6 up to 30 carbon atoms such as but not limited to phenyl, naphthyl, anthracenyl, phenanthracenyl, fluoranthenyl, chrysenyl, pyrenyl, biphenyl, terphenyl, picenyl, indenyl, biphenyl, indacenyl, benzocyclobutenyl, benzocyclooctenyl and the like, including fused benzo-C₄₋₈ cycloalkyl radicals (the latter being as defined above) such as, for instance, indanyl, tetrahydronaphthyl, fluorenyl and the like, all of the said radicals being optionally substituted with one or more substituents independently selected from the group consisting of halogen, amino, trifluoromethyl, hydroxyl, sulfhydryl and nitro, such as for instance 4-fluorophenyl, 4-chlorophenyl, 3,4-dichlorophenyl, 4-cyanophenyl, 2,6-dichlorophenyl, 2-fluorophenyl, 3-chlorophenyl, 3,5-dichlorophenyl and the like.

As used herein, e.g. with respect to a substituting radical such as the combination of substituents in positions 6 and 7 of the pteridine ring together with the carbon atoms in positions 6 and 7 of the pteridine ring, and unless otherwise stated, the term " homocyclic " means a mono- or polycyclic, saturated or mono-unsaturated or polyunsaturated hydrocarbon radical having from 4 up to 15 carbon atoms but including no heteroatom in the said ring; for instance said combination of substituents in positions 6 and 7 of the pteridine ring may form a C₂₋₆ alkylene radical, such as tetramethylene, which cyclizes with the carbon atoms in positions 6 and 7 of the pteridine ring.

As used herein with respect to a substituting radical (including the combination of substituents in positions 6 and 7 of the pteridine ring together with the carbon atoms in positions 6 and 7 of the pteridine ring), and unless otherwise stated, the term "heterocyclic" means a mono- or polycyclic, saturated or mono-unsaturated or polyunsaturated monovalent hydrocarbon radical having from 2 up to 15 carbon atoms and including one or more heteroatoms in one or more heterocyclic rings, each of said rings having from 3 to 10 atoms (and optionally further including one or more heteroatoms attached to one or more carbon atoms of said ring, for instance in the form of a carbonyl or thiocarbonyl or selenocarbonyl group, and/or to one or more heteroatoms of said ring, for instance in the form of a sulfone, sulfoxide, N-oxide, phosphate, phosphonate or selenium oxide group), each of said heteroatoms being independently selected from the group consisting of nitrogen, oxygen, sulfur, selenium and phosphorus, also including radicals wherein a heterocyclic ring is fused to one or more aromatic hydrocarbon rings for instance in the form of benzo-fused, dibenzo-fused and naphtho-fused heterocyclic radicals; within this definition are included heterocyclic radicals such as, but not limited to, diazepinyl, oxadiazinyl, thiadiazinyl, dithiazinyl, triazolonyl, diazepinonyl, triazepinyl, triazepinonyl, tetrazepinonyl, benzoquinolonyl, benzothiazinyl, benzothiazinonyl, benzoxa-thiiny, benzodioxinyl, benzodithiiny, benzoxazepinyl, benzothiazepinyl, benzodiazepinyl, benzodioxepinyl, benzodithiepinyl, benzoxazocinyl, benzothiazocinyl, benzodiazocinyl, benzoxathiocinyl, benzo-dioxocinyl, benzotrioxepinyl, benzoxathiazepinyl, benzoxadiazepinyl, benzothia-diazepinyl, benzotriazepinyl, benzoxathiepinyl, benzotriazinonyl, benzoxazolinonyl, azetidionyl, azaspirodecyl, dithiaspirodecyl, selenazinyl, selenazolyl, selenophenyl, hypoxanthinyl, azahypoxanthinyl, bipyrazinyl, bipyridinyl, oxazolidinyl, diselenopyrimidinyl, benzodioxocinyl, benzopyrenyl, benzopyranonyl, benzophenazinyl, benzoquinoliziny, dibenzocarbazoly, dibenzoacridinyl, dibenzophenazinyl, dibenzothiepinyl, dibenzooxepinyl, dibenzopyranonyl, dibenzoquinoxaliny, dibenzothiazepinyl, dibenzoiso-quinolonyl, tetraazaadamantyl, thiatetraazaadamantyl, oxauracil, oxazinyl, dibenzothiophenyl, dibenzofuranyl, oxazolonyl, azaindoly, azolonyl, thiazolonyl, thiazolonyl, thiazolidinyl, thiazanyl, pyrimidonyl, thiopyrimidonyl, thiamorpholinyl, azlactonyl, naphthindazolyl, naphthindolyl, naphthothiazolyl, naphthothioxolyl, naphthoxindolyl, naphthotriazolyl, naphtho-pyranyl, oxabicycloheptyl, azabenzimidazolyl, azacycloheptyl, azacyclooctyl, azacyclononyl, azabicyclononyl, tetrahydrofuryl, tetrahydropyranyl, tetrahydro-pyranyl, tetrahydroquinoleinyl, tetrahydrothienyl and dioxide thereof, dihydrothienyl dioxide, dioxindolyl, dioxinyl, dioxenyl, dioxazinyl, thioxanyl, thioxolyl, thiourazolyl, thiotriazolyl, thiopyranyl, thiopyronyl, coumarinyl, quinoleinyl, oxyquinoleinyl, quinuclidinyl, xanthinyl, dihydropyranyl, benzo-dihydrofuryl, benzothiopyronyl, benzothiopyranyl, benzoxazinyl, benzoxazolyl, benzodioxolyl, benzodioxanyl, benzothiadiazolyl, benzotriazinyl, benzo-thiazolyl, benzoxazolyl, phenothioxinyl, phenothiazolyl, phenothienyl (benzothiofuranyl), phenopyronyl, phenoxazolyl, pyridinyl, dihydropyridinyl, tetrahydropyridinyl, piperidinyl, morpholinyl, thiomorpholinyl, pyrazinyl,

pyrimidinyl, pyridazinyl, triazinyl, tetrazinyl, triazolyl, benzotriazolyl, tetrazolyl, imidazolyl, pyrazolyl, thiazolyl, thiadiazolyl, isothiazolyl, oxazolyl, oxadiazolyl, pyrrolyl, furyl, dihydrofuryl, furoyl, hydantoinyl, dioxolanyl, dioxolyl, dithianyl, dithienyl, dithiiny, thienyl, indolyl, indazolyl, benzofuryl, quinolyl, quinazolinyl, quinoxalyl, carbazolyl, phenoxazinyl, phenothiazinyl, xanthenyl, purinyl, benzothienyl, naphthothienyl, thianthrenyl, pyranyl, pyronyl, benzopyronyl, isobenzofuranyl, chromenyl, phenoxathiinyl, indoliziny, quinoliziny, isoquinolyl, phthalazinyl, naphthiridinyl, cinnoliny, pteridinyl, carboliny, acridiny, perimidiny, phenanthroliny, phenazinyl, phenothiazinyl, imidazoliny, imidazolidiny, benzimidazolyl, pyrazoliny, pyrazolidiny, pyrroliny, pyrrolidiny, piperazinyl, uridiny, thymidiny, cytidiny, aziriny, aziridinyl, diaziriny, diazolidiny, oxirany, oxaziridinyl, dioxirany, thiirany, azetyl, dihydroazetyl, azetidiny, oxetyl, oxetany, oxetanony, homopiperazinyl, homopiperidiny, thietyl, thietany, diazabicyclooctyl, diazetyl, diaziridinony, diaziridinethionyl, chromany, chromanony, thiochromany, thiochromanony, thiochromenyl, benzofurany, benzisothiazolyl, benzo-carbazolyl, benzochromony, benziso-alloxazinyl, benzocoumariny, thiocoumariny, phenometoxazinyl, phenoparoxazinyl, phentriazinyl, thiodiazinyl, thiodiazolyl, indoxyl, thioindoxyl, benzodiazinyl (e.g. phtalazinyl), phtalidyl, phtalimidiny, phtalazonyl, alloxazinyl, dibenzopyronyl (i.e. xanthony), xanthionyl, isatyl, isopyrazolyl, isopyrazolony, urazolyl, uraziny, uretiny, uretidiny, succiny, succinimido, benzylsultimyl, benzylsultamyl and the like, including all possible isomeric forms thereof, wherein each carbon atom of said heterocyclic ring may be independently substituted with a substituent selected from the group consisting of halogen, nitro, C₁₋₇ alkyl (optionally containing one or more functions or radicals selected from the group consisting of carbonyl (oxo), alcohol (hydroxyl), ether (alkoxy), acetal, amino, imino, oximino, alkylloximino, amino-acid, cyano, carboxylic acid ester or amide, nitro, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, C₁₋₇ alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkylamino, hydroxylalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl, sulfonamido and halogen), C₃₋₇ alkenyl, C₂₋₇ alkynyl, halo C₁₋₇ alkyl, C₃₋₁₀ cycloalkyl, aryl, arylalkyl, alkylaryl, alkylacyl, arylacyl, hydroxyl, amino, C₁₋₇ alkylamino, cycloalkylamino, alkenylamino, cyclo-alkenylamino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfhydryl, C₁₋₇ alkoxy, C₃₋₁₀ cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, thioaryl, thioheterocyclic, arylalkylthio, heterocyclic-substituted alkylthio, formyl, hydroxylamino, cyano, carboxylic acid or esters or thioesters or amides thereof, thiocarboxylic acid or esters or thioesters or amides thereof; depending upon the number of unsaturations in the 3 to 10 membered ring, heterocyclic radicals may be sub-divided into heteroaromatic (or "heteroaryl") radicals and non-aromatic heterocyclic radicals; when a heteroatom of the said

non-aromatic heterocyclic radical is nitrogen, the latter may be substituted with a substituent selected from the group consisting of C₁₋₇ alkyl, C₃₋₁₀ cycloalkyl, aryl, arylalkyl and alkylaryl.

As used herein with respect to a substituting radical, and unless otherwise stated, the terms "C₁₋₇ alkoxy", "C₃₋₁₀ cycloalkoxy", "aryloxy", "arylalkyloxy", "oxyheterocyclic",
5 "thio C₁₋₇ alkyl", "thio C₃₋₁₀ cycloalkyl", "arylthio", "arylalkylthio" and "thioheterocyclic" refer to substituents wherein a C₁₋₇ alkyl radical, respectively a C₃₋₁₀ cycloalkyl, aryl, arylalkyl or heterocyclic radical (each of them such as defined herein), are attached to an oxygen atom or a divalent sulfur atom through a single bond, such as but not limited to methoxy, ethoxy, propoxy, butoxy, pentoxy, isopropoxy, sec-butoxy, tert-butoxy, isopentoxy, cyclopropyloxy,
10 cyclobutyloxy, cyclopentyloxy, thiomethyl, thioethyl, thiopropyl, thiobutyl, thiopentyl, thiocyclopropyl, thiocyclobutyl, thiocyclopentyl, thiophenyl, phenyloxy, benzyloxy, mercaptobenzyl, cresoxy, and the like.

As used herein with respect to a substituting atom, and unless otherwise stated, the term halogen means any atom selected from the group consisting of fluorine, chlorine,
15 bromine and iodine.

As used herein with respect to a substituting radical, and unless otherwise stated, the term "halo C₁₋₇ alkyl" means a C₁₋₇ alkyl radical (such as above defined) in which one or more hydrogen atoms are independently replaced by one or more halogens (preferably fluorine, chlorine or bromine), such as but not limited to difluoromethyl, trifluoromethyl,
20 trifluoroethyl, octafluoropentyl, dodecafluoroheptyl, dichloromethyl and the like.

As used herein with respect to a substituting radical, and unless otherwise stated, the terms "C₂₋₇ alkenyl" designate a straight and branched acyclic hydrocarbon monovalent radical having one or more ethylenic unsaturations and having from 2 to 7 carbon atoms such as, for example, vinyl, 1-propenyl, 2-propenyl (allyl), 1-butenyl, 2-butenyl, 2-pentenyl, 3-
25 pentenyl, 3-methyl-2-butenyl, 3-hexenyl, 2-hexenyl, 2-heptenyl, 1,3-butadienyl, pentadienyl, hexadienyl, heptadienyl, heptatrienyl and the like, including all possible isomers thereof.

As used herein with respect to a substituting radical, and unless otherwise stated, the term "C₃₋₁₀ cycloalkenyl" mean a monocyclic mono- or polyunsaturated hydrocarbon monovalent radical having from 3 to 8 carbon atoms, such as for instance cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclopentadienyl, cyclohexenyl, cyclohexadienyl, cycloheptenyl,
30 cyclohepta-dienyl, cycloheptatrienyl, cyclooctenyl, cyclooctadienyl and the like, or a C₇₋₁₀ polycyclic mono- or polyunsaturated hydrocarbon mono-valent radical having from 7 to 10 carbon atoms such as dicyclopentadienyl, fencheryl (including all isomers thereof, such as α -pinolenyl), bicyclo[2.2.1]hept-2-enyl, bicyclo[2.2.1]hepta-2,5-dienyl, cyclo-fencheryl and the
35 like.

As used herein with respect to a substituting radical, and unless otherwise stated, the term "C₂₋₇ alkynyl" defines straight and branched chain hydrocarbon radicals containing one or more triple bonds and optionally at least one double bond and having from 2 to 7 carbon atoms such as, for example, acetylenyl, 1-propynyl, 2-propynyl, 1-butylenyl, 2-butylenyl, 2-

pentynyl, 1-pentynyl, 3-methyl-2-butynyl, 3-hexynyl, 2-hexynyl, 1-penten-4-ynyl, 3-penten-1-ynyl, 1,3-hexadien-1-ynyl and the like.

As used herein with respect to a substituting radical, and unless otherwise stated, the terms "arylalkyl", "arylalkenyl" and "heterocyclic-substituted alkyl" refer to an aliphatic saturated or ethylenically unsaturated hydrocarbon monovalent radical (preferably a C₁₋₇ alkyl or C₂₋₇ alkenyl radical such as defined above) onto which an aryl or heterocyclic radical (such as defined above) is already bonded, and wherein the said aliphatic radical and/or the said aryl or heterocyclic radical may be optionally substituted with one or more substituents independently selected from the group consisting of halogen, amino, hydroxyl, sulphydryl, C₁₋₇ alkyl, trifluoromethyl and nitro, such as but not limited to benzyl, 4-chlorobenzyl, 4-fluorobenzyl, 2-fluorobenzyl, 3,4-dichlorobenzyl, 2,6-dichlorobenzyl, 3-methylbenzyl, 4-methylbenzyl, 4-*ter*-butylbenzyl, phenylpropyl, 1-naphthylmethyl, phenylethyl, 1-amino-2-phenylethyl, 1-amino-2-[4-hydroxy-phenyl]ethyl, 1-amino-2-[indol-2-yl]ethyl, styryl, pyridylmethyl (including all isomers thereof), pyridylethyl, 2-(2-pyridyl)isopropyl, oxazolylbutyl, 2-thienylmethyl, pyrrolylethyl, morpholinylethyl, imidazol-1-yl-ethyl, benzodioxolylmethyl and 2-furylmethyl.

As used herein with respect to a substituting radical, and unless otherwise stated, the term "alkylaryl" and "alkyl-substituted heterocyclic" refer to an aryl or heterocyclic radical (such as defined above) onto which are bonded one or more aliphatic saturated or unsaturated hydrocarbon mono-valent radicals, preferably one or more C₁₋₇ alkyl, C₂₋₇ alkenyl or C₃₋₁₀ cycloalkyl radicals as defined above such as, but not limited to, o-toluyyl, m-toluyyl, p-toluyyl, 2,3-xylyl, 2,4-xylyl, 3,4-xylyl, o-cumenyl, m-cumenyl, p-cumenyl, o-cymenyl, m-cymenyl, p-cymenyl, mesityl, *ter*-butylphenyl, lutidinyl (i.e. dimethylpyridyl), 2-methylaziridinyl, methylbenzimidazolyl, methylbenzo-furanyl, methylbenzothiazolyl, methylbenzotriazolyl, methylbenzoxazolyl and methylbenzselenazolyl.

As used herein with respect to a substituting radical, and unless otherwise stated, the term "alkoxyaryl" refers to an aryl radical (such as defined above) onto which is (are) bonded one or more C₁₋₇ alkoxy radicals as defined above, preferably one or more methoxy radicals, such as, but not limited to, 2-methoxyphenyl, 3-methoxyphenyl, 4-methoxyphenyl, 3,4-dimethoxyphenyl, 2,4,6-trimethoxyphenyl, methoxynaphtyl and the like.

As used herein with respect to a substituting radical, and unless otherwise stated, the terms "alkylamino", "cycloalkylamino", "alkenyl-amino", "cycloalkenylamino", "arylamino", "arylalkylamino", "heterocyclic-substituted alkylamino", "heterocyclic-substituted arylamino", "heterocyclic amino", "hydroxyalkylamino", "mercaptoalkylamino" and "alkynylamino" mean that respectively one (thus monosubstituted amino) or even two (thus disubstituted amino) C₁₋₇ alkyl, C₃₋₁₀ cycloalkyl, C₂₋₇ alkenyl, C₃₋₁₀ cycloalkenyl, aryl, arylalkyl, heterocyclic-substituted alkyl, heterocyclic-substituted aryl, heterocyclic (provided in this case the nitrogen atom is attached to a carbon atom of the heterocyclic ring), mono- or polyhydroxy C₁₋₇ alkyl, mono- or polymercapto C₁₋₇ alkyl or C₂₋₇ alkynyl radical(s) (each of

them as defined herein, respectively) is/are attached to a nitrogen atom through a single bond such as but not limited to, anilino, benzylamino, methylamino, dimethylamino, ethylamino, diethylamino, isopropylamino, propenylamino, n-butylamino, ter-butylamino, dibutylamino, morpholinoalkylamino, 4-morpholinoanilino, hydroxymethylamino, β -hydroxyethylamino and ethynylamino; this definition also includes mixed disubstituted amino radicals wherein the nitrogen atom is attached to two such radicals belonging to two different sub-set of radicals, e.g. an alkyl radical and an alkenyl radical, or to two different radicals within the same sub-set of radicals, e.g. methylethylamino; among disubstituted amino radicals, symmetrically substituted are more easily accessible and thus usually preferred.

As used herein with respect to a substituting radical, and unless otherwise stated, the terms "(thio)carboxylic acid ester", "(thio)carboxylic acid thioester" and "(thio)carboxylic acid amide" refer to radicals wherein the carboxyl or thiocarboxyl group is directly attached to the pteridine ring (e.g. in the 6- and/or 7-position) and wherein said carboxyl or thiocarboxyl group is bonded to the hydrocarbonyl residue of an alcohol, a thiol, a polyol, a phenol, a thiophenol, a primary or secondary amine, a polyamine, an amino-alcohol or ammonia, the said hydrocarbonyl residue being selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, arylalkyl, alkylaryl, alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, arylamino, arylalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydroxyalkylamino, mercapto-alkylamino or alkynylamino (such as above defined, respectively).

As used herein with respect to a substituting radical, and unless otherwise stated, the term "amino-acid" refers to a radical derived from a molecule having the chemical formula $H_2N-CHR-COOH$, wherein R is the side group of atoms characterizing the amino-acid type; said molecule may be one of the 20 naturally-occurring amino-acids or any similar non naturally-occurring amino-acid.

As used herein and unless otherwise stated, the term "stereoisomer" refers to all possible different isomeric as well as conformational forms which the compounds of formula (I), (IV) and (V) may possess, in particular all possible stereochemically and conformationally isomeric forms, all diastereomers, enantiomers and/or conformers of the basic molecular structure. Some compounds of the present invention may exist in different tautomeric forms, all of the latter being included within the scope of the present invention.

As used herein and unless otherwise stated, the term "enantiomer" means each individual optically active form of a compound of the invention, having an optical purity or enantiomeric excess (as determined by methods standard in the art) of at least 80% (i.e. at least 90% of one enantiomer and at most 10% of the other enantiomer), preferably at least 90% and more preferably at least 98%.

As used herein and unless otherwise stated, the term "solvate" includes any combination which may be formed by a pteridine derivative of this invention with a suitable

inorganic solvent (e.g. hydrates) or organic solvent, such as but not limited to alcohols, ketones, esters and the like.

As used herein and unless otherwise stated, the terms " dihydro-pteridine derivative " and " tetrahydropteridine derivative " refer to the hydrogenation products of the pteridine derivatives having the general formula (I), i.e. derivatives wherein two hydrogen atoms are present in positions 5 and 6, or 7 and 8, of the pteridine ring, or wherein four hydrogen atoms are present in positions 5, 6, 7 and 8 of the said ring; such hydrogenated derivatives are easily accessible from the pteridine derivatives using hydrogenation methods well known in the art.

DETAILED DESCRIPTION OF THE INVENTION

An object of the invention is to provide a pharmaceutical composition having high immunosuppressive activity. Thus, the present invention relates in particular to the medical applications of a group of pteridine derivatives, their pharmaceutically acceptable salts, N-oxides, solvates, polymorphs, dihydro- and tetrahydroderivatives and enantiomers, possessing unexpectedly desirable pharmaceutical properties, in particular which are highly active immunosuppressive agents, and as such are useful in the treatment in transplant rejection and/or in the treatment of certain inflammatory diseases.

Surprisingly, the compounds of the present invention show a broader therapeutic spectrum profile than merely immunosuppressive activity, as is evidenced by the results obtained in the diversity of test procedures disclosed hereinbelow. A further advantageous feature of the compounds of the present invention resides in their excellent oral activity.

In the first embodiment of the invention, the novel pteridine derivatives are as defined in the general formula (I), wherein each of the substituents R₀, R₁, R₂, R₃, R₄ and R₅ may independently correspond to any of the definitions given above, in particular with any of the individual meanings (such as illustrated above) of generic terms used for substituting radicals such as, but not limited to, "C₁₋₇ alkyl ", " C₃₋₁₀ cycloalkyl ", " C₂₋₇ alkenyl ", " C₂₋₇ alkynyl ", " aryl ", " homocyclic ", " heterocyclic ", " halogen ", " C₃₋₁₀ cycloalkenyl ", "alkylaryl ", "arylalkyl ", "alkylamino", " cycloalkylamino ", "alkenylamino ", "alkynylamino", "arylamino", " arylalkylamino ", " heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, ", "hydroxyalkylamino ", " mercaptoalkylamino ", " alkynylamino ", " C₁₋₇ alkoxy", " C₃₋₁₀ cycloalkoxy ", "thio C₁₋₇ alkyl ", " thio C₃₋₁₀ cycloalkyl ", " halo C₁₋₇ alkyl ", "amino-acid " and the like.

In the second embodiment of the invention, the novel pteridine derivatives are as defined in the general formula (V), wherein each of the substituents R₂, R₆ and R₇ may independently correspond to any of the definitions given above, in particular with any of the individual meanings (such as illustrated above) of generic terms used for substituting radicals such as, but not limited to, " C₁₋₇ alkyl ", " C₂₋₇ alkenyl ", " C₂₋₇ alkynyl ", " aryl ", " homocyclic ", " heterocyclic ", " halogen ", "alkylaryl ", "arylalkyl ", "alkylamino", " cycloalkylamino ",

"alkenylamino ", "alkynylamino", "arylamino", " arylalkylamino ", " heterocyclic-substituted alkylamino ", " heterocyclic amino ", "heterocyclic-substituted arylamino ", " hydroxyalkylamino ", " mercaptoalkylamino ", " alkynylamino ", " C₁₋₇ alkoxy", " C₃₋₁₀ cycloalkoxy ", "thio C₁₋₇ alkyl ", " thio C₃₋₁₀ cycloalkyl ", " halo C₁₋₇ alkyl " and the like.

5

Stereoisomers of the compounds of this invention may be formed by using reactants in their single enantiomeric form wherever possible in the manufacturing process or by resolving the mixture of stereoisomers by conventional methods. One such method. is liquid chromatography using one or more suitable chiral stationary phases including, for example, poly-saccharides, in particular cellulose or amylose derivatives. Commercially available polysaccharide-based chiral stationary phases are ChiralCelTM CA, OA, OB, OC, OD, OF, 10 OG, OJ and OK, and ChiralpakTM AD, AS, OP(+) and OT(+). Appropriate eluents or mobile phases for use in combination with said polysaccharide-based chiral stationary phases are hydrocarbons such as hexane and the like, optionally admixed with an alcohol such as ethanol, isopropanol and the like. The above mixture of enantiomers may alternatively be 15 separated by making use of microbial resolution or by resolving the diastereoisomeric salts formed with chiral acids such as mandelic acid, camphorsulfonic acid, tartaric acid, lactic acid and the like or with chiral bases such as brucine and the like. The resolving agent may be cleaved from the separated diastereoisomers, e.g. by treatment with acids or bases, in order 20 to generate the pure enantiomers of the compounds of the invention. Conventional resolution methods were compiled e.g. by Jaques et al. in " Enantiomers, Racemates and Resolution " (Wiley Interscience, 1981).

In the general formula (II), the schematic notation (III)



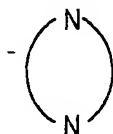
preferably means a heterocyclic group selected from the group consisting of:

- 25 - piperazin-1-yl,
 - homopiperazin-1-yl,
 - 4-imidazolin-1-yl,
 - imidazolidin-1-yl,
 - 2,3-dihydropyrazol-1-yl,
 30 - 2,3,4,5-tetrahydropyrazol-1-yl,
 - 3-pyrazolin-1-yl,
 - 4-pyrazolin-1-yl,
 - pyrazolidin-1-yl,
 - 2,3-dihydropyrazin-1-yl,
 35 - tetrahydropyrazin-1-yl,

- dihydropyrimidin-1-yl,
- tetrahydropyrimidin-1-yl,
- dihydropyridazin-1-yl,
- tetrahydropyridazin-1-yl,
- 5 - hexahydropyridazin-1-yl,
- dihydrofurazan-2-yl,
- tetrahydrofurazan-2-yl,
- dihydrophenazin-5-yl,
- dihydrotriazol-1-yl,
- 10 - dihydrotriazol-2-yl,
- tetrahydrotriazol-1-yl,
- tetrahydrotriazol-2-yl,
- dihydrotriazin-1-yl,
- tetrahydrotriazin-1-yl,
- 15 - tetrahydrooxadiazin-2-yl,
- tetrahydrothiadiazin-2-yl,
- dihydroindazol-1-yl,
- dihydroindazol-2-yl,
- tetrahydrophthalazin-2-yl,
- 20 - tetrahydrophthalazin-3-yl,
- tetrahydroquinoxalin-1-yl,
- tetrahydroquinazolin-1-yl,
- tetrahydroquinazolin-3-yl,
- dihydrocinnolin-1-yl,
- 25 - dihydrocinnolin-2-yl,
- tetrahydrocinnolin-1-yl,
- tetrahydrocinnolin-2-yl,
- dihydroperimidin-1-yl,
- tetrahydrodiazepin-1-yl, and
- 30 - oxides, sulfones and selenium oxides of the latter.

In the general formula (II), the said heterocyclic group may be substituted, at one or more carbon atoms, by a number n of substituents R_0 wherein n is an integer from 0 to 6 and wherein, when n is at least 2, each R_0 may be defined independently from the others. The presence of one or more such substituents R_0 is a suitable way for introducing chirality into the pteridine derivatives having the general formula (I) as well as into the polysubstituted 6-aminopyrimidines having the general formula (IV) and the polysubstituted 2,6-diaminopyrimidines having the general formula (VII). In practice, the choice of substituents R_0 may be restricted by the commercial availability of the substituted heterocyclic amine, depending upon the specific nature of the heterocyclic group.

More preferably the schematic notation (III)



represents a piperazin-1-yl group or a homopiperazin-1-yl group, in which case preferably n is 0, 1 or 2, and a representative example of the substituent R_0 is methyl or phenyl (such as for instance in 2-methylpiperazin-1-yl, 2-phenylpiperazin-1-yl and 2,5-dimethyl-piperazin-1-yl).

5 As shown in the general formula (II) taken together with the definition of R_1 , a requirement of an embodiment of the invention is that one of the two nitrogen atoms of the heterocyclic ring bears a substituent R_1 which has a carbonyl (oxo) or thiocarbonyl (thioxo) or sulfonyl function preferably immediately adjacent to the said nitrogen atom. In other words, this embodiment means that when R_1 is selected from, respectively, acyl, thioacyl, amide, 10 thioamide, sulfonyl, sulfinyl, carboxylate and thiocarboxylate, then R_1 together with the nitrogen atom to which it is attached forms, respectively, an amide, thioamide, urea, thiourea, sulfonamido, sulfinamido, carbamato or thiocarbamato group.

As already specified above, one or more of the substituents R_2 , R_3 , R_4 and R_5 of the pteridine ring may be a group represented by the general formula (II) with anyone of the 15 individual meanings of the substituent R_1 and anyone of the individual meanings of the optional substituent(s) R_0 . As will be apparent from the synthetic routes described hereinafter, preferably one or two of the substituents R_2 , R_3 , R_4 and R_5 of the pteridine ring are groups independently having said general formula (II). More preferably these substituents are R_4 and/or R_5 , i.e. the ones in positions 2 and/or 4 of the pteridine ring. When both positions 2 and 20 4 of the pteridine ring are substituted by groups (i.e. R_4 and R_5) having the general formula (II), these substituents may be the same (as shown in figure 8) or different (as shown in figures 6 and 7).

As already specified above, the remaining positions of the pteridine ring, i.e. the ones which are not substituted by a group represented by the general formula (II), may either be 25 unsubstituted (i.e. one, two or three of the relevant groups R_2 , R_3 , R_4 or R_5 is/are hydrogen atom) or be substituted independently from each other in the manner described hereinabove. Preferably one or two of said remaining positions, being more preferably selected from positions 2, 6 and 7, of the pteridine ring are substituted. When two such remaining positions (e.g. positions 2 and 6) of the pteridine ring are substituted in the manner described 30 hereinabove, the relevant substituents are preferably different from each other.

Some preferred pteridine derivatives having the general formula (I) according to the invention are more specifically illustrated in the following examples and defined in the following claims. For instance, useful pteridine species disclosed below include those wherein:

35 R_2 is selected from the group consisting of hydrogen, C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, preferably a phenyl group optionally substituted with one or more substituents selected from the group consisting of halogen, C_{1-7} alkyl and C_{1-7} alkoxy, more preferably

p-fluorophenyl, p-chlorophenyl, p-tolyl, p-acetamidophenyl or 3,4-dimethoxyphenyl, and/or

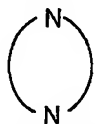
- R₃ is selected from the group consisting of hydrogen, C₁₋₇ alkyl, C₃₋₁₀ cycloalkyl, aryl and heteroaryl, preferably hydrogen, and/or
- 5 - R₄ is amino or a group represented by the general formula (II), and/or
- R₅ is a group selected from the group consisting of piperazin-1-yl, homopiperazin-1-yl, 2-methylpiperazin-1-yl, 2-phenylpiperazin-1-yl and 2,5-dimethylpiperazin-1-yl, the said group being substituted in the 4 position of the piperazinyl or homopiperazinyl ring, with a substituent R₁ which has a carbonyl (oxo) or thiocarbonyl (thioxo) or sulfonyl function (e.g.
- 10 R₁ is selected from the group consisting of acyl, thioacyl, amide, thioamide, sulfonyl, sulfinyl, carboxylate and thiocarboxylate) preferably immediately adjacent to the 4 position nitrogen atom of said ring.

Especially useful species of pteridine derivatives having the general formula (I) are those wherein one of the substituents R₄ and R₅ is a piperazin-1-yl group or a homopiperazin-1-yl group, said group being substituted in the 4 position with a substituent R₁, wherein R₁ is selected from the group consisting of:

- COR₈ wherein R₈ is selected from the group consisting of hydrogen; C₁₋₇ alkyl; C₃₋₁₀ cycloalkyl; aryl optionally substituted with one or more substituents selected from the group consisting of halogen, C₁₋₇ alkyl, cyano and C₁₋₇ alkoxy; heterocyclic optionally substituted with one or more halogen atoms; arylalkyl; aryloxyalkyl; arylalkoxyalkyl; alkoxyalkyl; arylalkoxy; aryloxy; arylalkenyl; heterocyclic-substituted alkyl; alkylamino and arylamino; representative but non limiting examples of R₈ are methyl, ethyl, pentyl, cyclohexyl, phenyl, 4-fluorophenyl, 4-chlorophenyl, 3,4-dichlorophenyl, 4-butylphenyl, 4-cyanophenyl, 2-methoxyphenyl, 3-methoxyphenyl, 4-pentoxypheyl, naphthyl, 2-thienyl, 4-pyridinyl, 1-tetrahydropyrrolyl, 2-tetrahydropyrrolyl, 2-furanyl, 3-furanyl, 2,4-dichloro-5-fluoro-3-pyridinyl, diethylamino, diisopropylamino, diphenylamino, phenyl-ethyl, 4-chlorobenzyl, phenoxyethyl, benzyloxyethyl, methoxyethyl, 2-thienylethyl, styryl, benzyloxy, phenoxy, 1-amino-2-phenylethyl, 1-amino-2-[4-hydroxyphenyl]ethyl and 1-amino-2-[indol-2-yl]ethyl; when R₄ is a N-formylpiperazin-1-yl group (i.e. R₈ is hydrogen), then preferably R₅ is not alkylamino, arylalkylamino or heterocyclic, or R₂ is not hydrogen, C₁₋₇ alkyl or aryl, or R₃ is not alkylamino, arylalkylamino or heterocyclic;
- CSR₉, wherein R₉ is selected from the group consisting of alkylamino and aryloxy, such as but not limited to dimethylamino and phenoxy;
- SO₂ R₁₀, wherein R₁₀ is selected from the group consisting of aryl and arylalkyl, such as but not limited to phenyl and benzyl; and
- 35 - R₁₁, wherein R₁₁ is selected from the group consisting of C₁₋₇ alkyl, aryl, arylalkyl, arylalkenyl, alkoxyalkyl, heterocyclic-substituted alkyl, cycloalkylalkyl, heterocyclic, C₃₋₁₀ cycloalkyl, alkylaminoalkyl, aryloxyalkyl, alkoxyaryl, ω-cyanoalkyl, ω-carboxylatoalkyl and carboxamidoalkyl.

The present invention further provides various processes and methods for making the novel pteridine derivatives having the general formula (I). As a general rule, the preparation of these compounds is based on the principle that, starting from a suitable pteridine precursor (a diaminopyrimidine), each of the substituents R_2 , R_3 , R_4 and R_5 may be introduced separately (except, of course, when R_2 together with R_3 forms a homocyclic or heterocyclic radical) without adversely influencing the presence of one or more substituents already introduced at other positions on the pteridine ring or the capacity to introduce further substituents later on.

Methods of manufacture have been developed by the present inventors which may be used alternatively to, or may be combined with, the methods of synthesis already known in the art of pteridine derivatives (depending upon the targeted final compound). For instance, methods for simultaneously introducing R_2 and R_3 in the form of a homocyclic or heterocyclic radical at positions 6 and 7 of the pteridine ring are already known from U.S. Patent No. 2,581,889. The synthesis of mono- and di-*N*-oxides of the pteridine derivatives of this invention can easily be achieved by treating the said derivatives with an oxidizing agent such as, but not limited to, hydrogen peroxide (e.g. in the presence of acetic acid) or a peracid such as chloroperbenzoic acid. Dihydro- and tetrahydropteridine derivatives of this invention can easily be obtained by catalytic hydrogenation of the corresponding pteridine derivatives, e.g. by placing the latter in a hydrogen atmosphere in the presence of platinum oxide or platinum. The methods for making the pteridine derivatives of the present invention will now be explained in more details by reference to the appended figures 1 to 9 wherein, unless otherwise stated hereinafter, each of the substituting groups or atoms R_2 , R_3 , R_4 and R_5 is as defined in formula (I) of the summary of the invention and, more specifically, may correspond to any of the individual meanings disclosed above. For a reason of convenience, each of figures 1 to 4 shows piperazin-1-yl as a representative example of the heterocyclic ring schematically represented as

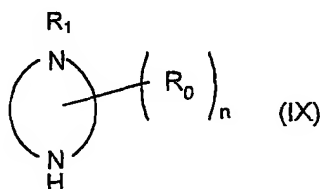


in the general formula (II), however it should be understood that the methods of the invention are not particularly limited to piperazin-1-yl but can be applied successfully to any other heterocyclic ring meeting the requirements specified hereinabove, in particular homopiperazin-1-yl.

In the description of the reaction steps involved in each figure, reference is made to the use of certain catalysts and/or certain types of solvents. It should be understood that each catalyst mentioned should be used in a catalytic amount well known to the skilled person with respect to the type of reaction involved. Solvents that may be used in the following reaction steps include various kinds of organic solvents such as protic solvents, polar aprotic solvents and non-polar solvents as well as aqueous solvents which are inert under the relevant reaction conditions. More specific examples include aromatic hydrocarbons, chlorinated

hydrocarbons, ethers, aliphatic hydrocarbons, alcohols, esters, ketones, amides, water or mixtures thereof, as well as supercritical solvents such as carbon dioxide (while performing the reaction under supercritical conditions). The suitable reaction temperature and pressure conditions applicable to each kind of reaction step will not be detailed herein but do not depart from the relevant conditions already known to the skilled person with respect to the type of reaction involved and the type of solvent used (in particular its boiling point).

Figure 1 schematically shows a first method for making trisubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II) and wherein R_2 or R_3 is hydrogen, as well as tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II). In step (a), a nitroso group is introduced on position 5 of the pyrimidine ring of a 2- R_4 -substituted 4-oxo-6-aminopyrimidine by using sodium nitrite under aqueous acidic conditions. Reduction of the nitroso group in step (b) is achieved either catalytically (Pt/H_2) in the presence of a protic solvent, or chemically using sodium dithionite or ammonium sulfide in water. Then in a next step (c), condensing the resulting 2- R_4 -substituted 4-oxo-5,6-diamino-pyrimidine with an α -ketoaldoxime bearing a radical R_2 , wherein R_2 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, in a protic solvent such as methanol under acidic conditions regioselectively yields a 4-oxopteridine bearing a R_4 substituent in position 2 and a R_2 substituent in position 6 of the pteridine ring. Alternatively, a 2- R_4 -substituted 4-oxo-7- R_3 -substituted pteridine derivative can be obtained in step (d) by reacting the 2- R_4 -substituted 4-oxo-5,6-diamino-pyrimidine with a monosubstituted glyoxal bearing the group R_3 , wherein R_3 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, under neutral or basic conditions. Alternatively, a 2- R_4 -substituted-4-oxo-6,7-disubstituted pteridine derivative can be obtained in step (e) by reacting the 2- R_4 -substituted 4-oxo-5,6-diamino-pyrimidine with a disubstituted glyoxal bearing groups R_2 and R_3 , wherein each of R_2 and R_3 is independently selected (i.e. R_2 and R_3 may be identical or different) from the group consisting of C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, under neutral or basic conditions. Activation of the (tautomeric) hydroxyl substituent in position 4 of the pteridine ring for a nucleophilic displacement reaction occurs in step (f) by preparing the corresponding 4-[(1,2,4)-triazolyl] pteridine derivative, e.g. using $POCl_3$ or 4-chlorophenyl phosphorodichloridate and 1,2,4-triazole in pyridine as solvent. When R_4 is an amino group, protection of R_4 may further be necessary before carrying out this reaction. The amino group can be protected for instance by an acetyl group, which can be hydrolysed back to the amino group in a next step. Nucleophilic substitution is performed in step (g) by mixing the triazolyl pteridine derivative with a nucleophile having the general formula (IX) :



wherein R_0 and n are as already defined above with respect to formula (II) and wherein R_1 is hydrogen or is as defined above with respect to formula (II), such as, but not limited to, piperazine or an appropriate N-alkylpiperazine, N-aryl piperazine or N-alkylaryl piperazine, at room temperature in a polar aprotic solvent such as 1,4-dioxane. When piperazine is introduced in step (g), then in step (h), the second nitrogen atom of the piperazin-1-yl substituent in position 4 of the pteridine ring can be coupled with the desired carboxylic acid or thio-carboxylic acid chloride or sulfonyl chloride $R_1\text{Cl}$ at room temperature in a solvent such as pyridine.

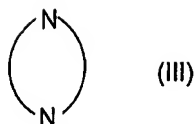
Representative but non limiting examples of commercially available N-alkylpiperazines, N-aryl piperazines and N-alkylaryl piperazines that can suitably be used in this method, as well as in some of the further methods described herein, include 1-cyclohexylpiperazine, 1-cyclopentylpiperazine, 1-(2,6-dichlorobenzyl)-piperazine, 1-(3,4-dichlorophenyl)-piperazine, 1-[2-(dimethylamino)-ethyl]-piperazine, 1-[3-(dimethylamino)-propyl]-piperazine, 1-(3,4-dimethylphenyl)piperazine, 1-(2-ethoxyethyl)-piperazine, 1-isobutylpiperazine, 1-(1-methyl-piperidin-4-yl-methyl)-piperazine, 1-(2-nitro-4-trifluoromethylphenyl)-piperazine, 1-(2-phenoxyethyl)-piperazine, 1-(1-phenylethyl)-piperazine, 2-(piperazin-1-yl)-acetic acid ethyl ester, 2-(piperazin-1-yl)-acetic acid *N*-methyl-*N*-phenyl amide, 2-(piperazin-1-yl)-acetic acid *N*-(2-thiazolyl)-amide, 2-[2-(piperazin-1-yl)-ethyl]-1,3-dioxolan-3-(1-piperazinyl)propionitrile, 1-[(2-pyridyl)-methyl]piperazine and 1-thiazol-2-yl-piperazine.

Figure 2 schematically shows a second method for making trisubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II) and wherein R_2 or R_3 is hydrogen, as well as tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II). In step (a), the diazonium salt of *p*-chloroaniline is first formed by using sodium nitrite under aqueous acidic conditions and then reacted with a 2- R_4 -substituted 4-chloro-6-amino-pyrimidine to yield an azo intermediate. In step (b), the chlorine atom in position 4 of the pyrimidinyl ring is replaced by nucleophile having the above general formula (IX) such as, but not limited to, a piperazinyl group or an appropriate N-alkylpiperazinyl, N-aryl piperazinyl or N-alkylaryl piperazinyl group. Reductive cleavage of the azo compound then yields the corresponding 2- R_4 -substituted 4-(piperazin-1-yl)-5,6-diaminopyrimidine in step (c). Condensation of the latter with an α -ketoaldehyde bearing a radical R_2 , wherein R_2 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, in a protic solvent such as methanol under acidic conditions leads in step (d) to the formation of a 2- R_4 -substituted 4-(piperazin-1-yl)-6- R_2 -substituted pteridine. Alternatively, a 2- R_4 -substituted 4-(piperazin-1-yl)-7- R_3 -substituted pteridine derivative can be obtained in step (e) by reacting the 2- R_4 -substituted 4-(piperazin-1-yl)-5,6-diaminopyrimidine with a monosubstituted glyoxal bearing the group R_3 , wherein R_3 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, under neutral or basic conditions. Alternatively, a 2- R_4 -substituted-4-(piperazin-1-yl)-6,7-disubstituted pteridine derivative can be obtained in step (f) by reacting the 2- R_4 -substituted 4-(piperazin-1-yl)-5,6-diamino-pyrimidine with a disubstituted glyoxal bearing

groups R_2 and R_3 , wherein each of R_2 and R_3 is independently selected (i.e. R_2 and R_3 may be identical or different) from the group consisting of C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, under neutral or basic conditions. When a piperazinyl group is introduced at position 4 of the pyrimidine scaffold in step (b), then coupling of the second nitrogen atom of the piperazin-1-yl group with an acid or sulfonyl chloride R_1Cl can occur in the last step (g) in the same way as in the first method above.

Figure 3 schematically shows a third method for making trisubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II) and wherein R_2 or R_3 is hydrogen, as well as tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II). This method starts from the pyrimidine derivative obtained after step (b) of the first method above. Formation of the pteridine ring occurs in step (b) through reaction of the said pyrimidine derivative with a suitable α -ketoaldoxime bearing a radical R_2 , wherein R_2 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, in a protic solvent such as methanol under acidic conditions. Alternatively, a 2- R_4 -substituted 4-oxo-7- R_3 -substituted pteridine derivative can be obtained in step (c) by reacting the 2- R_4 -substituted 4-oxo-5,6-diaminopyrimidine with a monosubstituted glyoxal bearing the group R_3 , wherein R_3 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, under neutral or basic conditions. Alternatively, a 2- R_4 -substituted 4-oxo-6,7-disubstituted pteridine derivative can be obtained in step (d) by reacting the 2- R_4 -substituted 4-oxo-5,6-diamino-pyrimidine with a disubstituted glyoxal bearing groups R_2 and R_3 , wherein each of R_2 and R_3 is independently selected (i.e. R_2 and R_3 may be identical or different) from the group consisting of C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, under neutral or basic conditions. A nucleophilic group such as, but not limited to, piperazin-1-yl or N-alkylpiperazinyl, N-arylpiperazinyl or N-alkylarylpiperazinyl is then directly introduced, in step (e), at position 4 of the pteridine ring by reaction of the 2- R_4 -substituted 4-oxopteridine with a nucleophile having the above general formula (IX) such as, but not limited to, piperazine or an appropriate N-alkylpiperazine, N-arylpiperazine or N-alkylarylpiperazine, and 1,1,1,3,3,3-hexamethyldisilazane as a reagent. When piperazine was used in step (e), then in step (f) coupling of the second nitrogen atom of the piperazin-1-yl group with an acid or sulfonyl chloride R_1Cl can proceed in the same way as in the first method above.

Figure 4 schematically shows a fourth method for making trisubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II) and wherein R_2 or R_3 is hydrogen, as well as tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II). In this method, R_4 preferably is amino. In a first step (a), the chlorine atom at position 4 of the pyrimidine ring of a 2- R_4 -substituted 4-chloro-6-aminopyrimidine is displaced by an appropriate R_1 -N-monosubstituted piperazine wherein R_1 may be acyl, alkyl, aryl, alkylaryl or sulfonyl, thus e.g. yielding a novel polysubstituted 2,6-diaminopyrimidine having the general formula (VII) wherein $n = 0$, R_7 is hydrogen and the heterocyclic ring having the formula (III)



is for instance a piperazinyl group.

Many *N*-monosubstituted piperazines required for step (a) are commercially available, such as for instance:

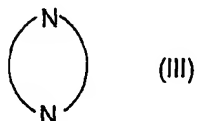
- 5 - 1-(2-furoyl) piperazine,
- 1-(4-chlorobenzenesulfonyl) piperazine ,
- 1-(4-fluorobenzoyl) piperazine,
- 1-(4-methoxyphenylsulfonyl) piperazine,
- 1-(ethoxycarbonyl) piperazine,
- 10 - 1-(tetrahydro-2-furoyl) piperazine,
- 1-(thien-2-ylcarbonyl) piperazine,
- 1-[(2,3-dihydro-1,4-benzodioxin-2-yl)carbonyl] piperazine,
- 1-acetyl piperazine,
- 1-formyl piperazine, and
- 15 - 1-methanesulfonyl piperazine.

N-acyl-, *N*-thioacyl- or *N*-sulfonyl- monosubstituted piperazines which are not commercially available may easily be prepared by reacting piperazine with any commercially available carboxylic acid chloride, thiocarboxylic acid chloride or sulfonyl chloride under standard acylation, thioacylation or sulfonylation conditions .

- 20 The method shown in figure 4 is also applicable while starting, in the first step (a), from an appropriate R_1 -monosubstituted homopiperazine wherein R_1 may be acyl or sulfonyl, thus e.g. yielding a novel polysubstituted 2,6-diaminopyrimidine having the general formula (VII) wherein $n = 0$, R_7 is hydrogen and the heterocyclic ring having the formula (III) is a homopiperazinyl group. Commercially available monosubstituted homopiperazines required
- 25 for such step (a) are, for instance, *N*-acetylhomopiperazine, 1-[3-chloro-5-(trifluoromethyl)-2-pyridyl]-homopiperazine, 1-[4-(trifluoromethyl)pyrimidin-2-yl]-homopiperazine, 1-(4-fluorobenzyl)-homopiperazine, 1-(2-chloro-6-fluoro-benzyl)-homopiperazine, 1-(5-nitro-2-pyridyl)-homopiperazine and (5-isoquinoline-sulfonyl)-homopiperazine. *N*-acyl-, *N*-thioacyl- or *N*-sulfonyl- monosubstituted homopiperazines which are not commercially available may easily
- 30 be prepared by reacting homopiperazine with any commercially available carboxylic acid chloride, thiocarboxylic acid chloride or sulfonyl chloride under standard acylation, thioacylation or sulfonylation conditions.

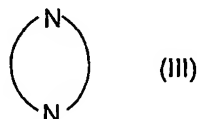
Introduction of a nitroso group at position 5 of the pyrimidine ring occurs in step (b) under aqueous acidic conditions in the presence of sodium nitrite, thus e.g. yielding a novel

polysubstituted 2,6-diaminopyrimidine having the general formula (VII) wherein $n = 0$, R_7 is nitroso and the heterocyclic ring having the formula (III):



is for instance a piperazinyl group (as shown in figure 4) or a homopiperazinyl group (not shown in figure 4).

- 5 Reduction of the nitroso functionality of this intermediate into a free amino group is then effected in step (c) by means of reducing agents such as $\text{Na}_2\text{S}_2\text{O}_4$ or $(\text{NH}_4)_2\text{S}$ in water, or catalytically (Pt/H_2) in the presence of a protic solvent, thus e.g. yielding a novel polysubstituted 2,5,6-triaminopyrimidine having the general formula (VII) wherein $n = 0$, R_7 is amino and the heterocyclic ring having the general formula (III)



- 10 is for instance a piperazinyl group (as shown in figure 4) or a homopiperazinyl group (not shown in figure 4).

- In order to regioselectively obtain a 2,4,6-trisubstituted pteridine derivative, the substituted 5,6-diaminopyrimidine is then reacted in step (d) with an α -ketoaldoxime bearing the group R_2 , wherein R_2 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, under acidic conditions in the presence of a solvent such a methanol. Alternatively, a 2,4,7-trisubstituted pteridine derivative can be obtained in step (f) by reacting the substituted 5,6-diaminopyrimidine with a monosubstituted glyoxal bearing the group R_3 , wherein R_3 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, under neutral or basic conditions. Alternatively, a 2,4,6,7-tetrasubstituted pteridine derivative can be obtained in step (e) by reacting the substituted 5,6-diaminopyrimidine with a disubstituted glyoxal bearing the groups R_2 and R_3 , wherein each of R_2 and R_3 is independently selected (i.e. R_2 and R_3 may be identical or different) from the group consisting of C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, under neutral or basic conditions.

- Figure 5 schematically shows a fifth method for making trisubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II) and wherein R_3 is hydrogen. In particular, figure 5 shows a scheme making use of a coupling compound such as a carboxylic or sulfonic acid having the general formula $R_{11}\text{ZOOH}$, wherein Z is selected from the group consisting of the carbon atom and the SO group, and wherein R_{11} is a group selected from the group consisting of C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl, heteroaryl and alkylaryl, wherein the said group R_{11} is optionally substituted with one or more substituents selected from the group consisting of halogen, amino and protected amino groups, and wherein the said substituent may be at any position (such as the α position or the ω position) with respect

to the carboxylic or sulfonic acid group. In particular, the said coupling compound may be a carboxylic acid $R_{11}CO_2H$ or a sulfonic acid $R_{11}SO_3H$ or a preferably amino-protected naturally-occurring amino-acid (e.g. L-alanine, L-phenyl-alanine, L-tyrosine, L-proline, L-tryptophane, L-leucine, L-isoleucine, L-lysine, L-valine, glycine, L-histidine, L-serine, L-arginine, L-aspartic acid, L-cysteine or L-glutamine) or synthetic non naturally-occurring amino-acid. This method provides coupling, in one single step (a), of the said compound to the second nitrogen atom of a heterocyclic ring having the general formula (III) and being substituted at position 4 of a pteridine ring. That is, this method starts for instance from a pteridine intermediate such as obtained after step (g) of the scheme shown in figure 1, or after any of steps (d), (e) and (f) of the scheme shown in figure 2, or after step (e) of the scheme shown in figure 3. When the coupling compound is a carboxylic acid $R_{11}CO_2H$, the said pteridine intermediate is reacted with the coupling compound preferably in an aprotic solvent, such as dimethyl-formamide or dichloromethane or mixtures thereof, and in the presence of a suitable coupling reagent, such as 1,3-dicyclohexylcarbodiimide or 1,3-diisopropylcarbodiimide or diisopropylethyl-amine or o-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium tetrafluoroborate. As is usual for this kind of coupling reaction, all free amino groups of the amino-acids are most preferably protected before carrying out the coupling reaction. When the amino-acid has two or more amino groups, the protecting groups for the said amino groups may be the same or different. A few examples of amino-protecting groups are benzyloxycarbonyl (which may be introduced by reaction of the desired amino-acid with benzylchloroformate under alkaline conditions, e.g. making use of sodium hydroxide or hydrogenocarbonate) and 9-fluorenylmethoxycarbonyl (which may be introduced by reaction of the desired amino-acid with 9-fluorenylmethyl chloroformate). Another example of an amino-protecting group is a tert-butoxycarbonyl group which may be introduced by reaction of the desired amino-acid with di-tert-butyl dicarbonate under alkaline conditions. Other suitable amino-protecting groups include triphenylmethyl (trityl) and trifluoroacetyl groups. First, the amino-protected amino-acid is coupled to the second nitrogen atom of the heterocyclic ring e.g. by using any method conventional in peptide synthesis. Finally, in order to afford for instance the desired (4-substituted-piperazin-1-yl) or (4-substituted-homopiperazin-1-yl) pteridine derivative, the amino-protecting group is removed by deprotection methods conventional in the art, such as:

- when the amino-protecting group is a phenylmethoxycarbonyl group, cleavage of the benzylic ether function by hydrogenolysis, e.g. using H_2 , Pd-C at about 25°C, or under strongly acidic conditions (e.g. making use of bromhydric acid), or
- when the amino-protecting group is a tert-butoxycarbonyl group, by treatment with an acid, e.g. using aqueous hydrochloric acid or trifluoroacetic acid, under conditions mild enough to avoid further cleavage of the molecule, or
- when the amino-protecting group is a 9-fluorenylmethoxycarbonyl group, by treatment with a base such as piperidine.

Figure 6 schematically shows a sixth method for making trisubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II) and wherein R_3 or R_2 is hydrogen, as well as tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II). In step (a), the thiol function of 2-mercapto-4,6-diaminopyrimidine is alkylated, preferably methylated by reaction with methyl iodide in the presence of a solvent such as ethanol, in order to yield 2-thiomethyl-4,6-diaminopyrimidine. Introduction of a nitroso group in the 5-position of the pyrimidine ring is then achieved in step (b) by using sodium nitrite under aqueous acidic conditions. In step (c), the methylthio group in the 2-position is exchanged for a group R_4 by reaction with an appropriate nucleophile, wherein R_4 is as defined above and preferably is primary or secondary amino, C_{1-7} alkoxy, aryloxy, C_{3-10} cycloalkoxy, heteroaryloxy, mercapto C_{1-7} alkyl, mercaptoaryl, mercapto C_{3-10} cycloalkyl or mercapto-heteroaryl. Reduction of the nitroso group is then achieved in step (d) either catalytically (Pt/H_2) in the presence of a protic solvent or chemically using sodium dithionite or ammonium sulfide in the presence of water. Then in step (e), the resulting 2- R_4 -substituted-4,5,6-triaminopyrimidine is condensed, under acidic conditions in the presence of a solvent such as methanol, with an α -ketoaldoxime bearing the group R_2 , wherein R_2 may be C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, into a 2,6-substituted-4-aminopteridine derivative. Alternatively, the corresponding 2,7-substituted-4-aminopteridine derivative can be obtained in step (f) by reacting the 2- R_4 -substituted-4,5,6-triaminopyrimidine with a monosubstituted glyoxal bearing a group R_3 , wherein R_3 may be inter alia C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl. Alternatively, a 2- R_4 -substituted 4-amino-6,7-disubstituted pteridine derivative can be obtained in step (g) by reacting the 2- R_4 -substituted 4,5,6-triaminopyrimidine with a disubstituted glyoxal bearing groups R_2 and R_3 , wherein each of R_2 and R_3 is independently selected (i.e. R_2 and R_3 may be identical or different) from the group consisting of C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, under neutral or basic conditions. In step (h), acidic or basic hydrolysis of the amino group at position 4 of the pteridine ring is performed and results in the corresponding 4-oxopteridine derivative. In step (i), the hydroxyl group of the tautomeric form of the latter is activated by nucleophilic displacement, e.g. by preparing the 4-[(1,2,4)-triazolyl] pteridine derivative. Finally in a first part of step (j), a nucleophilic displacement is performed by mixing the said 4-triazolylpteridine derivative with a nucleophile having the above general formula (IX).

When this nucleophile, and optionally also the nucleophile used in step (c), has a heterocyclic ring containing at least two nitrogen atoms, the second nitrogen atom of each heterocyclic ring can be acylated, thioacylated or sulfonylated in a last part of step (j) by treating the intermediate with an appropriate carboxylic acid chloride, thiocarboxylic acid chloride or sulfonyl chloride R_1Cl in an aprotic solvent such as dimethyl-formamide, pyridine or dichloromethane and, if necessary, in the presence of a base such as a tertiary amine (e.g. triethylamine).

Figure 7 schematically shows a seventh method for making trisubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II) and wherein R_3 or R_2 is hydrogen, as well as tetrasubstituted pteridine derivatives having the formula (I) wherein R_5 is a group having the formula (II), starting from the 2-thiomethyl-5-nitroso-4,6-diaminopyrimidine obtained after step (b) of the scheme shown in figure 6. Reduction of the nitroso group is achieved in step (a) either catalytically (Pt/H_2) in the presence of a protic solvent or chemically using sodium dithionite or ammonium sulfide in the presence of water. Then in step (b), 2-thiomethyl-4,5,6-triaminopyrimidine is condensed, under acidic conditions in the presence of a solvent such as methanol, with an α -ketoaldoxime bearing the group R_2 , wherein R_2 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl, thus regioselectively yielding a 2-thiomethyl-4-amino-6- R_2 -substituted-pteridine derivative. Alternatively, the corresponding 2-thiomethyl-4-amino-7- R_3 -substituted pteridine is obtained in step (c) by reacting 2-thiomethyl-4,5,6-triamino-pyrimidine with a monosubstituted glyoxal bearing a group R_3 , wherein R_3 may be *inter alia* C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl or heteroaryl. Alternatively, the corresponding 2-thiomethyl-4-amino-6- R_2 -7- R_3 -substituted pteridine is obtained in step (d) by reacting 2-thiomethyl-4,5,6-triamino-pyrimidine with a disubstituted glyoxal bearing groups R_2 and R_3 , wherein each of R_2 and R_3 is independently selected (i.e. R_2 and R_3 may be identical or different) from the group consisting of C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, under neutral or basic conditions. In step (e), the methylthio group in the 2-position is oxidized to the corresponding sulfone by using oxidizing agents such as chloroperoxybenzoic acid in chloroform or hydrogen peroxide in acetic acid. The methylsulfonyl group is easily exchanged in step (f) by reaction with a nucleophile having the general formula (IX) above. When this nucleophile has an heterocyclic ring containing at least two nitrogen atoms, the second nitrogen atom of said heterocyclic ring can be acylated, thioacylated or sulfonylated in step (f) by treatment with an appropriate carboxylic acid chloride, thiocarboxylic acid chloride or sulfonyl chloride R_1Cl in a aprotic solvent such as dimethylformamide, pyridine or dichloromethane and, if necessary, in the presence of a base such as a tertiary amine (e.g. triethylamine). In step (g), acidic or basic hydrolysis of the amino group at position 4 of the pteridine ring is performed and results in the corresponding 4-oxopteridine derivative. In step (h), the hydroxyl group of the tautomeric form of the latter is activated by nucleophilic displacement, e.g. by preparing the 4-[(1,2,4)-triazolyl] pteridine derivative. Finally in a first part of step (i), a nucleophilic displacement is performed by mixing the said 4-triazolylpteridine derivative with a nucleophile having the general formula (IX) above. When this nucleophile has a heterocyclic ring containing at least two nitrogen atoms, the second nitrogen atom of each heterocyclic ring can be acylated or sulfonylated in a last part of step (i) by treating the intermediate with an appropriate acid chloride or sulfonyl chloride R_1Cl in a aprotic solvent such as dimethylformamide, pyridine or dichloromethane and, if necessary, in the presence of a base such as a tertiary amine (e.g. triethylamine).

Figure 8 schematically shows a method for making trisubstituted (wherein R₂ or R₃ is hydrogen) and tetrasubstituted pteridine derivatives having the formula (I) wherein R₄ and R₅ are identical groups having the formula (II). In step (a), a nitro group is introduced in position 5 of a 6-amino-2,4-dioxypyrimidine under strongly acidic conditions (e.g. HNO₃, H₂SO₄). Then, in step (b), both hydroxyl groups from the tautomeric form are converted to chloro groups by treatment with a chlorinating agent such as POCl₃ or SOCl₂. Both chloro groups are then displaced in step (c) with a nucleophile having the above general formula (IX), thus yielding novel 6-aminopyrimidines having the general formula (IV) wherein R₁ is hydrogen and R₆ is nitro. The nitro group of the latter is then reduced in step (d) to an amino group by treatment with a reducing agent (e.g. Pt/H₂), thus yielding novel 6-aminopyrimidines having the general formula (IV) wherein R₁ is hydrogen and R₆ is amino. Finally, reaction of the latter with an α -ketoaldoxime bearing the group R₂, wherein R₂ may be *inter alia* aryl, C₁₋₇ alkyl, C₁₋₁₀ cycloalkyl or heteroaryl, regioselectively yields the desired 2,4,6-trisubstituted pteridine derivative in step (e). Alternatively, reaction of the 2,4-substituted-5,6-diaminopyrimidine from step (d) with a monosubstituted glyoxal bearing a group R₃ wherein R₃ may be *inter alia* C₁₋₇ alkyl, C₃₋₁₀ cycloalkyl, aryl or heteroaryl yields the desired 2,4,7-trisubstituted pteridine derivative in step (f). Alternatively, reaction of the 2,4-substituted-5,6-diaminopyrimidine from step (d) with a disubstituted glyoxal bearing groups R₂ and R₃, wherein each of R₂ and R₃ is independently selected (i.e. R₂ and R₃ may be identical or different) from the group consisting of C₁₋₇ alkyl, C₃₋₁₀ cycloalkyl, aryl and heteroaryl, under neutral or basic conditions, yields the desired 2,4,6,7-tetrasubstituted pteridine derivative in step (g). Finally, when the nucleophile used in step (c) has a heterocyclic ring containing at least two nitrogen atoms, the second nitrogen atom can be acylated, thioacylated or sulfonylated in a last step (not shown in the figure) by treatment with an appropriate carboxylic acid chloride, thiocarboxylic acid chloride or sulfonyl chloride R₁Cl in a aprotic solvent such as dimethylformamide, pyridine or dichloromethane and, if necessary, in the presence of a base such as a tertiary amine (e.g. triethylamine).

Figure 9 schematically shows a method for making trisubstituted and tetrasubstituted pteridine derivatives having the formula (I) wherein R₅ is a piperazine substituted in position 4 with a ureido group. In the first step (a), a *N*-alkyl substituted aniline derivative is reacted with the commercially available *N,N*-carbonyldiimidazole using a polar aprotic solvent such as tetrahydrofuran or 1,4-dioxane. The reactivity of the carbamoyl imidazole was then increased by *N*-alkylation of the imidazole moiety in step (b) by reaction of *N*-alkyl aniline carbamoyl imidazole with an appropriate alkyl halide, such as, for example methyl iodide, leading to the formation of the corresponding imidazolium salt. Subsequent addition, in step (c), of a 4-*N*-piperazinopteridine (optionally substituted in positions 2 and/or 6 and/or 7) in an aprotic solvent such as dimethylformamide, pyridine or dichloromethane and, if necessary, in the presence of a base such as a tertiary amine (e.g. triethylamine) yields the desired substituted pteridine derivative.

Especially useful species of pteridine derivatives having the general formula (V) are those wherein:

- R₂ is 2-phenylethylamino, 2-thienylmethylamino, pyrrolidino, benzylamino or piperidino, and/or
- 5 - R₆ is aryl such as phenyl or 3,4-dimethoxyphenyl, and/or
- R₇ is hydrogen.

The preparation of such compounds is extensively described in some of the following examples, as well as in the above description of figures 6 and 7, wherein it can be seen that they are able to serve as intermediates for making pteridine derivatives having the general
10 formula (I).

In another particular embodiment, the invention relates to a group of pteridine derivatives, as well as pharmaceutical compositions comprising such pteridine derivatives as active principle, having the above general formula (I) or the general formula (V) and being in the form of a pharmaceutically acceptable salt. The latter include any therapeutically active
15 non-toxic addition salt which compounds having the general formula (I) or the general formula (V) are able to form with a salt-forming agent. Such addition salts may conveniently be obtained by treating the pteridine derivatives of the invention with an appropriate salt-forming acid or base. For instance, pteridine derivatives having basic properties may be converted into the corresponding therapeutically active, non-toxic acid addition salt form by treating the free
20 base form with a suitable amount of an appropriate acid following conventional procedures. Examples of such appropriate salt-forming acids include, for instance, inorganic acids resulting in forming salts such as but not limited to hydrohalides (e.g. hydrochloride and hydrobromide), sulfate, nitrate, phosphate, diphosphate, carbonate, bicarbonate, and the like; and organic monocarboxylic or dicarboxylic acids resulting in forming salts such as, for
25 example, acetate, propanoate, hydroxyacetate, 2-hydroxypropanoate, 2-oxopropanoate, lactate, pyruvate, oxalate, malonate, succinate, maleate, fumarate, malate, tartrate, citrate, methanesulfonate, ethanesulfonate, benzoate, 2-hydroxybenzoate, 4-amino-2-hydroxybenzoate, benzene-sulfonate, p-toluenesulfonate, salicylate, p-aminosalicylate, pamoate, bitartrate, camphorsulfonate, edetate, 1,2-ethanedisulfonate, fumarate,
30 glucoheptonate, gluconate, glutamate, hexylresorcinate, hydroxynaphtoate, hydroxyethanesulfonate, mandelate, methylsulfate, pantothenate, stearate, as well as salts derived from ethanedioic, propanedioic, butanedioic, (Z)-2-butenedioic, (E)-2-butenedioic, 2-hydroxybutanedioic, 2,3-dihydroxybutane-dioic, 2-hydroxy-1,2,3-propanetricarboxylic and cyclohexanesulfamic acids and the like.

35 Pteridine derivatives of the general formula (I) or (V) having acidic properties may be converted in a similar manner into the corresponding therapeutically active, non-toxic base addition salt form. Examples of appropriate salt-forming bases include, for instance, inorganic bases like metallic hydroxides such as but not limited to those of alkali and alkaline-earth metals like calcium, lithium, magnesium, potassium and sodium, or zinc, resulting in the

corresponding metal salt; organic bases such as but not limited to ammonia, alkylamines, benzathine, hydrabamine, arginine, lysine, N,N'-dibenzylethylenediamine, chloroprocaine, choline, diethanolamine, ethylene-diamine, N-methylglucamine, procaine and the like.

Reaction conditions for treating the pteridine derivatives having the general formula (I) or (V) of this invention with an appropriate salt-forming acid or base are similar to standard conditions involving the same acid or base but different organic compounds with basic or acidic properties, respectively. Preferably, in view of its use in a pharmaceutical composition or in the manufacture of medicament for treating specific diseases, the pharmaceutically acceptable salt will be designed, i.e. the salt-forming acid or base will be selected so as to impart greater water-solubility, lower toxicity, greater stability and/or slower dissolution rate to the pteridine derivative of this invention.

The present invention further provides the use of a pteridine derivative represented by the general formula (I) or the general formula (V), or a pharmaceutically acceptable salt or a solvate thereof, as a biologically-active ingredient, i.e. an active principle, especially as a medicine or a diagnostic agent or for the manufacture of a medicament or a diagnostic kit. In particular the said medicament may be for the prevention or treatment of a pathologic condition selected from the group consisting of:

- immune disorders, in particular organ and cells transplant rejections, and autoimmune disorders,
- cardiovascular disorders,
- allergic conditions,
- disorders of the central nervous system,
- TNF- α -related disorders,
- viral diseases, and
- cell proliferative disorders.

The pathologic conditions and disorders concerned by the said use, and the corresponding methods of prevention or treatment, are detailed hereinbelow. Any of the uses mentioned with respect to the present invention may be restricted to a non-medical use (e.g. in a cosmetic composition), a non-therapeutic use, a non-diagnostic use, a non-human use (e.g. in a veterinary composition), or exclusively an *in-vitro* use, or a use with cells remote from an animal.

The invention further relates to a pharmaceutical composition comprising:

- (a) one or more pteridine derivatives represented by the general formula (I) or the general formula (V), and
- (b) one or more pharmaceutically acceptable carriers.

In a third embodiment, this invention provides combinations, preferably synergistic combinations, of one or more pteridine derivative represented by the general formula (I) or the general formula (V) with one or more biologically-active drugs being preferably selected from the group consisting of immunosuppressant and/or immunomodulator drugs, antineoplastic

drugs, anti-histamines, inhibitors of allergy-causative agents (anti-allergic drugs) and antiviral agents. As is conventional in the art, the evaluation of a synergistic effect in a drug combination may be made by analyzing the quantification of the interactions between individual drugs, using the median effect principle described by Chou et al. in *Adv. Enzyme*
 5 *Reg.* (1984) 22:27. Briefly, this principle states that interactions (synergism, additivity, antagonism) between two drugs can be quantified using the combination index (hereinafter referred as CI) defined by the following equation:

$$CI_x = \frac{ED_x^{1c}}{ED_x^{1a}} + \frac{ED_x^{2c}}{ED_x^{2a}}$$

wherein ED_x is the dose of the first or respectively second drug used alone (1a, 2a), or in
 10 combination with the second or respectively first drug (1c, 2c), which is needed to produce a given effect. The said first and second drug have synergistic or additive or antagonistic effects depending upon $CI < 1$, $CI = 1$, or $CI > 1$, respectively. As will be explained in more detail herein-below, this principle may be applied to a number of desirable effects such as, but not limited to, an activity against transplant rejection, an activity against immunosuppression or
 15 immunomodulation, an activity against allergy or an activity against cell proliferation.

For instance the present invention relates to a pharmaceutical composition or combined preparation having synergistic effects against immuno-suppression or immunomodulation and containing:

- (a) one or more immunosuppressant and/or immunomodulator drugs, and
- 20 (b) at least one pteridine derivative represented by the general formula (I) or the general formula (V), and
- (c) optionally one or more pharmaceutical excipients or pharmaceutically acceptable carriers, for simultaneous, separate or sequential use in the treatment or prevention of autoimmune disorders and/or in transplant-rejections.

25 Suitable immunosuppressant drugs for inclusion in the synergistic compositions or combined preparations of this invention belong to a well known therapeutic class. They are preferably selected from the group consisting of cyclosporin A, substituted xanthines (e.g. methylxanthines such as pentoxifylline), daltroban, sirolimus, tacrolimus, rapamycin (and derivatives thereof such as defined below), leflunomide (or its main active metabolite
 30 A771726, or analogs thereof called malononitrilamides), mycophenolic acid and salts thereof (including the sodium salt marketed under the trade name Mofetil[®]), adrenocortical steroids, azathioprine, brequinar, gusperimus, 6-mercaptopurine, mizoribine, chloroquine, hydroxychloroquine and monoclonal antibodies with immunosuppressive properties (e.g. etanercept, infliximab or kineret). Adrenocortical steroids within the meaning of this invention
 35 mainly include glucocorticoids such as but not limited to ciprocinonide, desoxycorticosterone, fludrocortisone, flumoxonide, hydrocortisone, naflocort, procinonide, timobesone, tipredane, dexamethasone, methylprednisolone, methotrexate, prednisone, prednisolone, triamcinolone and pharmaceutically acceptable salts thereof. Rapamycin derivatives as referred herein

include O-alkylated derivatives, particularly 9-deoxorapamycins, 26-dihydrorapamycins, 40-O-substituted rapamycins and 28,40-O,O-disubstituted rapamycins (as disclosed in U.S. Patent No. 5,665,772) such as 40-O-(2-hydroxy) ethyl rapamycin – also known as SDZ-RAD -,
pegylated rapamycin (as disclosed in U.S. Patent No. 5,780,462), ethers of 7-desmethylrapamycin (as disclosed in U.S. Patent No. 6,440,991) and polyethylene glycol
5 esters of SDZ-RAD (as disclosed in U.S. Patent No. 6,331,547).

Suitable immunomodulator drugs for inclusion into the synergistic immunomodulating pharmaceutical compositions or combined preparations of this invention are preferably selected from the group consisting of acemannan, amiprilose, bucillamine, dimepranol,
10 ditiocarb sodium, imiquimod, Inosine Pranobex, interferon- β , interferon- γ , lentinan, levamisole, lisophylline, pidotimod, romurtide, platonin, procodazole, propagermanium, thymomodulin, thymopentin and ubenimex.

Synergistic activity of the pharmaceutical compositions or combined preparations of this invention against immunosuppression or immuno-modulation may be readily
15 determined by means of one or more lymphocyte activation tests. Usually activation is measured via lymphocyte proliferation. Inhibition of proliferation thus always means immunosuppression under the experimental conditions applied. There exist different stimuli for lymphocyte activation, in particular:

- a) co-culture of lymphocytes of different species (mixed lymphocyte reaction, hereinafter referred as MLR) in a so-called mixed lymphocyte culture test: lymphocytes expressing different minor and major antigens of the HLA-DR type (= alloantigens) activate each other non-specifically;
- b) a CD3 assay wherein there is an activation of the T-lymphocytes via an exogenously added antibody (OKT3). This antibody reacts against a CD3 molecule located on the
25 lymphocyte membrane which has a co-stimulatory function. Interaction between OKT3 and CD3 results in T-cell activation which proceeds via the Ca^{2+} /calmodulin/calcineurin system and can be inhibited e.g. by cyclosporin A (hereinafter referred as CyA);
- c) a CD28 assay wherein specific activation of the T-lymphocyte proceeds via an exogenously added antibody against a CD28 molecule which is also located on the
30 lymphocyte membrane and delivers strong co-stimulatory signals. This activation is Ca^{2+} -independent and thus cannot be inhibited by CyA.

Determination of the immunosuppressing or immunomodulating activity of the pteridine derivatives of this invention, as well as synergistic combinations comprising them, is preferably based on the determination of one or more, preferably at least three lymphocyte
35 activation *in vitro* tests, more preferably including at least one of the MLR test, CD3 assay and CD28 assay referred above. Preferably the lymphocyte activation *in vitro* tests used include at least two assays for two different clusters of differentiation preferably belonging to the same general type of such clusters and more preferably belonging to type I transmembrane proteins. Optionally the determination of the immuno-suppressing or immunomodulating

activity may be performed on the basis of other lymphocyte activation *in vitro* tests, for instance by performing a TNF- α assay or an IL-1 assay or an IL-6 assay or an IL-10 assay or an IL-12 assay or an assay for a cluster of differentiation belonging to a further general type of such clusters and more preferably belonging to type II transmembrane proteins such as, but
5 not limited to, CD69, CD 71 or CD134.

The synergistic effect may be evaluated by the median effect analysis method described herein-before. Such tests may for instance, according to standard practice in the art, involve the use of equipment, such as flow cytometer, being able to separate and sort a number of cell subcategories at the end of the analysis, before these purified batches can be
10 analysed further.

Synergistic activity of the pharmaceutical compositions of this invention in the prevention or treatment of transplant rejection may be readily determined by means of one or more leukocyte activation tests performed in a Whole Blood Assay (hereinafter referred as WBA) described for instance by Lin et al. in *Transplantation* (1997) 63:1734-1738. WBA used
15 herein is a lymphoproliferation assay performed *in vitro* using lymphocytes present in the whole blood, taken from animals that were previously given the pteridine derivative, and optionally the other immunosuppressant drug, *in vivo*. Hence this assay reflects the *in vivo* effect of substances as assessed by an *in vitro* read-out assay. The synergistic effect may be evaluated by the median effect analysis method described herein-before. Various organ
20 transplantation models in animals are also available *in vivo*, which are strongly influenced by different immunogenicities, depending on the donor and recipient species used and depending on the nature of the transplanted organ. The survival time of transplanted organs can thus be used to measure the suppression of the immune response.

The pharmaceutical composition or combined preparation with synergistic activity against
25 immunosuppression or immunomodulation according to this invention may contain the pteridine derivative of formula (I) or the general formula (V) over a broad content range depending on the contemplated use and the expected effect of the preparation. Generally, the pteridine derivative content of the combined preparation is within the range of 0.1 to 99.9% by weight, preferably from 1 to 99% by weight, more preferably from 5 to 95% by weight.

30 The invention further relates to a composition or combined preparation having synergistic effects against cell proliferation and containing:

- (a) one or more antineoplastic drugs, and
- (b) at least one pteridine derivative represented by the general formula (I) or the general formula (V), and
- 35 (c) optionally one or more pharmaceutical excipients or pharmaceutically acceptable carriers, for simultaneous, separate or sequential use in the treatment or prevention of cell proliferative disorders.

- Suitable antineoplastic drugs for inclusion into the synergistic antiproliferative pharmaceutical compositions or combined preparations of this invention are preferably selected from the group consisting of alkaloids, alkylating agents (including but not limited to alkyl sulfonates, aziridines, ethylenimines, methylmelamines, nitrogen mustards and nitrosoureas), antibiotics, antimetabolites (including but not limited to folic acid analogs, purine analogs and pyrimidine analogs), enzymes, interferon and platinum complexes. More specific examples include acivicin; aclarubicin; acodazole; acronine; adozelesin; aldesleukin; altretamine; ambomycin; ametantrone; aminoglutethimide; amsacrine; anastrozole; anthramycin; asparaginase; asperlin; azacitidine; azetepa; azotomycin; batimastat; benzodepa; bicalutamide; bisantrene; bisnafide; bizelesin; bleomycin; brequinar; bropirimine; busulfan; cactinomycin; calusterone; caracemide; carbetimer; carboplatin; carmustine; carubicin; carzelesin; cedefingol; chlorambucil; cirolemycin; cisplatin; cladribine; crisnatol; cyclophosphamide; cytarabine; dacarbazine; dactinomycin; daunorubicin; decitabine; dexornaplatin; dezaguanine; diaziquone; docetaxel; doxorubicin; droloxifene; dromostanolone; duazomycin; edatrexate; eflomithine; elsamitrucin; enloplatin; enpromate; epipropidine; epirubicin; erbulozole; esorubicin; estramustine; etanidazole; ethiodized oil I 131; etoposide; etoprine; fadrozole; fazarabine; fenretinide; floxuridine; fludarabine; fluorouracil; flurocitabine; fosquidone; fostriecin; gemcitabine; Gold 198; hydroxyurea; idarubicin; ifosfamide; ilmofofosine; interferon α -2a; interferon α -2b; interferon α -n1; interferon α -n3; interferon β -1a; interferon γ -1b; iroplatin; irinotecan; lanreotide; letrozole; leuprolide; liarozole; lometrexol; lomustine; losoxantrone; masoprocol; maytansine; mechlorethamine; megestrol; melengestrol; melphalan; menogaril; mercaptopurine; methotrexate; metoprine; meturedopa; mitindomide; mitocarcin; mitocromin; mitogillin; mitomalcin; mitomycin; mitosper; mitotane; mitoxantrone; mycophenolic acid; nocodazole; nogala-mycin; ormaplatin; oxisuran; paclitaxel; pegaspargase; peliomycin; pentamustine; peplomycin; perfosfamide; pipobroman; piposulfan; piroxantrone; plicamycin; plomestane; porfimer; porfiromycin; prednimustine; procarbazine; puromycin; pyrazofurin; riboprine; rogletimide; safingol; semustine; simtrazene; sparfosate; sparsomycin; spirogermanium; spiromustine; spiroplatin; streptonigrin; streptozocin; strontium 89 chloride; sulofenur; talisomycin; taxane; taxoid; tecogalan; tegafur; teloxantrone; temoporfin; teniposide; teroxirone; testolactone; thiamiprine; thioguanine; thiotepa; tiazofurin; tirapazamine; topotecan; toremifene; trestolone; tricinbine; trimetrexate; triptorelin; tubulozole; uracil mustard; uredepa; vapreotide; verteporfin; vinblastine; vincristine; vindesine; vinepidine; vinylicinate; vinleurosine; vinorelbine; vinrosidine; vinzolidine; vorozole; zeniplatin; zinostatin; zorubicin; and their pharmaceutically acceptable salts.
- Other suitable anti-neoplastic compounds include 20-epi-1,25 dihydroxyvitamin D₃; 5-ethynyluracil; abiraterone; aclarubicin; acylfulvene; adecypenol; adozelesin; aldesleukin; ALL-TK antagonists; altretamine; ambamustine; amidox; amifostine; aminolevulinic acid; amrubicin; amsacrine; anagrelide; anastrozole; andrographolide; angiogenesis inhibitors; antagonist D; antagonist G; antarelix; anti-dorsalizing morphogenetic protein-1; anti-

androgens such as, but not limited to, benorterone, cioteronel, cyproterone, delmadinone, oxendolone, topterone, zanoterone and their pharmaceutically acceptable salts; anti-estrogens such as, but not limited to, clometherone; delmadinone; nafoxidine; nitromifene; raloxifene; tamoxifen; toremifene; trioxifene and their pharmaceutically acceptable salts;

5 antineoplaston; antisense oligonucleotides; aphidicolin glycinate; apoptosis gene modulators; apoptosis regulators; apurinic acid; ara-CDP-DL-PTBA; arginine deaminase; asulacrine; atamestane; atrimustine; axinastatin; azasetron; azatoxin; azatyrosine; baccatin III derivatives; balanol; batimastat; BCR/ABL antagonists; benzochlorins; benzoylstauosporine; β -lactam derivatives; β -alethine; betaclamycin B; betulinic acid; bFGF inhibitor; bicalutamide;

10 bisantrene; bisazirindinylspermine; bisnafide; bistratene A; bizelesin; breflate; bropirimine; budotitane; buthionine sulfoximine; calcipotriol; calphostin C; camptothecin derivatives; canarypox IL-2; capecitabine; carboxamide-amino-triazole; carboxyamidotriazole; CaRest M3; CARN 700; cartilage derived inhibitor; carzelesin; casein kinase inhibitors; castanospermine; cecropin B; cetorelix; chlorins; chloroquinoxaline sulfonamide; cicaprost; cis-porphyrin;

15 clomifene and analogues thereof; clotrimazole; collismycin A and B; combretastatin and analogues thereof; conagenin; crambescidin 816; cryptophycin and derivatives thereof; curacin A; cyclopentanthraquinones; cycloplatam; cypemycin; cytarabine; cytolytic factor; cytostatin; dacliximab; dehydroidemnin B; deslorelin; dexifosfamide; dexrazoxane; dexverapamil; didemnin B; didox; diethylnorspermine; dihydro-5-azacytidine; dihydrotaxol;

20 dioxamycin; diphenyl spiromustine; docosanol; dolasetron; doxifluridine; droloxifene; dronabinol; duocamycin SA; ebselen; ecomustine; edelfosine; edrecolomab; elemene; emitefur; epristeride; estrogen agonists and antagonists; exemestane; filgrastim; finasteride; flavopiridol; flezelastine; fluasterone; fluorodaunorunicin; forfenimex; formestane; fotemustine; gadolinium texaphyrin; gallium nitrate; galocitabine; ganirelix; gelatinase inhibitors; glutathione

25 inhibitors; hepsulfam; heregulin; hexamethylene bisacetamide; hypericin; ibandronic acid; idoxifene; idramantone; ilomastat; imidazoacridones; imiquimod; immunostimulant peptides; insulin-like growth factor-1 receptor inhibitor; interferon agonists; iobenguane; iododoxorubicin; ipomeanol; irinotecan; iroplact; irsogladine; isobengazole; isohomohalicondrin B; itasetron; jasplakinolide; kahalalide F; lamellarin-N; leinamycin;

30 lenograstim; lentinan; leptolstatin; leukemia inhibiting factor; leuprorelin; levamisole; liarozole; lissoclinamide; lobaplatin; lombricine; lonidamine; lovastatin; loxoribine; lurtotecan; lutetium texaphyrin; lysofylline; mannostatin A; marimastat; masoprocol; maspin; matrilysin inhibitors; matrix metalloproteinase inhibitors; merbarone; meterelin; methioninase; metoclopramide; MIF inhibitors; mifepristone; miltefosine; mirimostim; mitoguazone; mitolactol; mitonafide;

35 mitotoxin fibroblast growth factor-saporin; mofarotene; molgramostim; human chorionic gonadotrophin monoclonal antibody; mopidamol; mycaperoxide B; myriaporone; N-acetyldinaline; N-substituted benzamides; nafarelin; nagrestip; naloxone; pentazocine; napavin; naphterpin; nartograstim; nedaplatin; nemorubicin; neridronic acid; neutral endopeptidase; nilutamide; nisamycin; nitric oxide modulators; nitroxide antioxidant; nitrullyn;

octreotide; okicenone; onapristone; ondansetron; ondansetron; oracin; osaterone; oxaliplatin; oxaunomycin; palauamine; palmitoylrhizoxin; pamidronic acid; panaxytriol; panomifene; parabactin; pazelliptine; peldesine; pentosan; pentostatin; pentozole; perflubron; perillyl alcohol; phenazinomycin; phenylacetate; phosphatase inhibitors; picibanil; pilocarpine; 5 pirarubicin; piritrexim; placetin A and B; plasminogen activator inhibitor; propyl bis-acridone; prostaglandin J2; proteasome inhibitors; protein kinase C inhibitors; protein tyrosine phosphatase inhibitors; purine nucleoside phosphorylase inhibitors; purpurins; pyrazoloacridine; raltitrexed; ramosetron; ras farnesyl protein transferase inhibitors; ras inhibitors; ras-GAP inhibitors; retelliptine; rhenium 186 etidronate; rhizoxin; retinamide; 10 rohitukine; romurtide; roquinimex; rubiginone B1; ruboxyl; saintopin; sarcophytol A; sargramostim; sizofiran; sobuzoxane; sodium borocaptate; sodium phenylacetate; solverol; somatomedin binding protein; sonermin; sparfosic acid; spicamycin D; splenopentin; spongistatin 1; squalamine; stem-cell division inhibitors; stipiamide; stromelysin inhibitors; sulfinosine; suradista; suramin; swainsonine; tallimustine; tamoxifen; tauromustine; 15 tazarotene; tecogalan; tellurapyrylium; telomerase inhibitors; temozolomide; tetrachlorodecaoxide; tetrazomine; thaliblastine; thiocoraline; thrombopoietin; thymalfasin; thymopoietin receptor agonist; thymotrnan; thyroid stimulating hormone; tin ethyl etiopurpurin; titanocene; topsentin; tretinoin; triacetyluridine; tropisetron; turosteride; tyrosine kinase inhibitors; tyrphostins; ubenimex; urogenital sinus-derived growth inhibitory factor; urokinase 20 receptor antagonists; variolin B; velaresol; veramine; verdins; verteporfin; vinxaltine; vitaxin; zanoterone; zilascorb; and their pharmaceutically acceptable salts.

The compounds of this invention may also be administered in combination with anti-cancer agents which act by arresting cells in the G2-M phases due to stabilized microtubules. In addition to Taxol (paclitaxel), and analogs and derivatives thereof, other examples of anti- 25 cancer agents which act by this mechanism include without limitation the following marketed drugs and drugs in development: erbulozole, dolastatin, mivobulin isethionate, discodermolide, altorhyrtins, spongistatins, cemadotin hydrochloride, epothilones desoxyepothilone, 16-aza-epothilone, 21-aminoepothilone, 21-hydroxyepothilone, 26-fluoroepothilone, auristatin, soblidotin, cryptophycin, vitilevuamide, tubulysin, canadensol, 30 centaureidin, oncocidin, fijianolide, laulimalide, narcosine, nascapine, hemiasterlin, vanadocene acetylacetonate, monsatrol, inanocine, eleutherobins, caribaeoside, caribaeolin, halichondrin, diazonamide, taccalonolide, diozostatin, phenylahistin, myoseverin, resverastatin phosphate sodium, and their pharmaceutically acceptable salts.

Synergistic activity of the pharmaceutical compositions or combined preparations of this 35 invention against cell proliferation may be readily determined by means of one or more tests such as, but not limited to, the measurement of the radioactivity resulting from the incorporation of ³H-thymidine in culture of tumor cell lines. For instance, different tumor cell lines are selected in order to evaluate the anti-tumor effects of the test compounds, such as but not limited to:

- RPMI1788: human Peripheral Blood Leucocytes (PBL) Caucasian tumor line,
- Jurkat: human acute T cell leukemia,
- EL4: C57Bl/6 mouse lymphoma, or
- THP-1: human monocyte tumor line.

5 Depending on the selected tumor cell line, different culture media may be used, such as for example:

- for RPMI1788 and THP-1: RPMI-1640 + 10% FCS + 1% NEAA + 1% sodium pyruvate + 5×10^{-5} mercapto-ethanol + antibiotics (G-418 0.45 µg/ml).
- for Jurkat and EL4: RPMI-1640 + 10% FCS + antibiotics (G-418 0.45 µg/ml).

10 In a specific embodiment of the synergy determination test, the tumor cell lines are harvested and a suspension of 0.27×10^6 cells/ml in whole medium is prepared. The suspensions (150 µl) are added to a microtiter plate in triplicate. Either complete medium (controls) or the test compounds at the test concentrations (50 µl) are added to the cell suspension in the microtiter plate. The cells are incubated at 37°C under 5% CO₂ for about 16
15 hours. ³H-thymidine is added, and the cells incubated for another 8 hours. The cells are harvested and radioactivity is measured in counts per minute (CPM) in a β-counter. The ³H-thymidine cell content, and thus the measured radioactivity, is proportional to the proliferation of the cell lines. The synergistic effect is evaluated by the median effect analysis method as disclosed herein-before.

20 The pharmaceutical composition or combined preparation with synergistic activity against cell proliferation according to this invention may contain the pteridine derivative of the general formula (I) or the general formula (V) over a broad content range depending on the contemplated use and the expected effect of the preparation. Generally, the pteridine derivative content of the combined preparation is within the range of 0.1 to 99.9% by weight, preferably from 1 to 99% by weight, more preferably from 5 to 95% by weight.
25

The invention further relates to a pharmaceutical composition or combined preparation having synergistic effects against a viral infection and containing:

- (a) one or more anti-viral agents, and
- (b) at least one pteridine derivative represented by the general formula (I) or the general
30 formula (V), and
- (c) optionally one or more pharmaceutical excipients or pharmaceutically acceptable carriers, for simultaneous, separate or sequential use in the treatment or prevention of a viral infection.

Suitable anti-viral agents for inclusion into the synergistic antiviral compositions or combined preparations of this invention include, for instance, retroviral enzyme inhibitors
35 belonging to categories well known in the art, such as HIV-1 IN inhibitors, nucleoside reverse transcriptase inhibitors (e.g. zidovudine, lamivudine, didanosine, stavudine, zalcitabine and the like), non-nucleoside reverse transcriptase inhibitors (e.g. nevirapine, delavirdine and the like), other reverse transcriptase inhibitors (e.g. foscarnet sodium and the like), and HIV-1 protease inhibitors (e.g. saquinavir, ritonavir, indinavir, nelfinavir and the like). Other suitable

antiviral agents include for instance acemannan, acyclovir, adefovir, alovudine, alvircept, amantadine, aranotin, arildone, atevirdine, avridine, cidofovir, cipamfylline, cytarabine, desciclovir, disoxaril, edoxudine, enviroxime, famciclovir, famotone, fiacitabine, fialuridine, floxuridine, fosarilate, fosfonet, ganciclovir, idoxuridine, kethoxal, lobucavir, memotone, methisazone, penciclovir, pirodavis, somantadine, sorivudine, tilorone, trifluridine, valaciclovir, vidarabine, viroxime, zinviroxime, moroxydine, podophyllotoxin, ribavirine, rimantadine, stallimycine, statolon, tromantadine and xenazoic acid, and their pharmaceutically acceptable salts.

Especially relevant to this aspect of the invention is the inhibition of the replication of viruses selected from the group consisting of picorna-, toga-, bunya, orthomyxo-, paramyxo-, rhabdo-, retro-, arena-, hepatitis B-, hepatitis C-, hepatitis D-, adeno-, vaccinia-, papilloma-, herpes-, corona-, varicella- and zoster-virus, in particular human immunodeficiency virus (HIV). Synergistic activity of the pharmaceutical compositions or combined preparations of this invention against viral infection may be readily determined by means of one or more tests such as, but not limited to, the isobologram method, as previously described by Elion et al. in *J. Biol. Chem.* (1954) 208:477-488 and by Baba et al. in *Antimicrob. Agents Chemother.* (1984) 25:515-517, using EC_{50} for calculating the fractional inhibitory concentration (hereinafter referred as FIC). When the minimum FIC index corresponding to the FIC of combined compounds (e.g., $FIC_x + FIC_y$) is equal to 1.0, the combination is said to be additive; when it is between 1.0 and 0.5, the combination is defined as subsynergistic, and when it is lower than 0.5, the combination is by defined as synergistic. When the minimum FIC index is between 1.0 and 2.0, the combination is defined as subantagonistic and, when it is higher than 2.0, the combination is defined as antagonistic.

The pharmaceutical composition or combined preparation with synergistic activity against viral infection according to this invention may contain the pteridine derivative of the general formula (I) or the general formula (V) over a broad content range depending on the contemplated use and the expected effect of the preparation. Generally, the pteridine derivative content of the combined preparation is within the range of 0.1 to 99.9% by weight, preferably from 1 to 99% by weight, more preferably from 5 to 95% by weight.

The pharmaceutical compositions and combined preparations according to this invention may be administered orally or in any other suitable fashion. Oral administration is preferred and the preparation may have the form of a tablet, aqueous dispersion, dispersable powder or granule, emulsion, hard or soft capsule, syrup, elixir or gel. The dosing forms may be prepared using any method known in the art for manufacturing these pharmaceutical compositions and may comprise as additives sweeteners, flavoring agents, coloring agents, preservatives and the like. Carrier materials and excipients are detailed hereinbelow and may include, *inter alia*, calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and disintegrating agents, binding agents and the like. The pharmaceutical composition or combined preparation of this invention may be included in a

gelatin capsule mixed with any inert solid diluent or carrier material, or has the form of a soft gelatin capsule, in which the ingredient is mixed with a water or oil medium. Aqueous dispersions may comprise the biologically active composition or combined preparation in combination with a suspending agent, dispersing agent or wetting agent. Oil dispersions may
5 comprise suspending agents such as a vegetable oil. Rectal administration is also applicable, for instance in the form of suppositories or gels. Injection (e.g. intramuscularly or intraperitoneously) is also applicable as a mode of administration, for instance in the form of injectable solutions or dispersions, depending upon the disorder to be treated and the condition of the patient.

10 Auto-immune disorders to be prevented or treated by the pharmaceutical compositions or combined preparations of this invention include both systemic auto-immune diseases such as but not limited to lupus erythematosus, psoriasis, vasculitis, polymyositis, scleroderma, multiple sclerosis, ankylosing spondylitis, rheumatoid arthritis and Sjögren syndrome; auto-immune endocrine disorders such as thyroiditis; and organ-specific auto-
15 immune diseases such as but not limited to Addison disease, hemolytic or pernicious anemia, Goodpasture syndrome, Graves disease, idiopathic thrombocytopenic purpura, insulin-dependent diabetes mellitus, juvenile diabetes, uveitis, Crohn's disease, ulcerative colitis, pemphigus, atopic dermatitis, autoimmune hepatitis, primary biliary cirrhosis, autoimmune pneumonitis, autoimmune carditis, myasthenia gravis, glomerulonephritis and spontaneous
20 infertility.

Transplant rejections to be prevented or treated by the pharmaceutical compositions or combined preparations of this invention include the rejection of transplanted or grafted organs or cells (both allografts and xenografts), such as but not limited to host versus graft reaction disease. The term "organ" as used herein means all organs or parts of organs in mammals,
25 in particular humans, such as but not limited to kidney, lung, bone marrow, hair, cornea, eye (vitreous), heart, heart valve, liver, pancreas, blood vessel, skin, muscle, bone, intestine or stomach. "Rejection" as used herein mean all reactions of the recipient body or of the transplanted organ which in the end lead to cell or tissue death in the transplanted organ or adversely affect the functional ability and viability of the transplanted organ or the recipient. In
30 particular, this means acute and chronic rejection reactions. Also included in this invention is preventing or treating the rejection of cell transplants and xenotransplantation. The major hurdle for xenotransplantation is that even before the T lymphocytes, responsible for the rejection of allografts, are activated, the innate immune system, especially T-independent B lymphocytes and macrophages are activated. This provokes two types of severe and early
35 acute rejection called hyper-acute rejection and vascular rejection, respectively. The present invention addresses the problem that conventional immunosuppressant drugs like cyclosporin A are ineffective in xeno-transplantation. The ability of the compounds of this invention to suppress T-independent xeno-antibody production as well as macrophage activation may be

evaluated in the ability to prevent xenograft rejection in athymic, T-deficient mice receiving xenogenic hamster-heart grafts.

Cell proliferative disorders to be prevented or treated by the pharmaceutical compositions or combined preparations of this invention include any kind of tumor progression or invasion or metastasis inhibition of a cancer, preferably one selected from the group consisting of lung cancer, leukaemia, ovarian cancer, sarcoma, Kaposi's sarcoma, meningioma, colon cancer, lymph node tumor, glioblastoma multiforme, prostate cancer or skin carcinose.

CNS disorders to be prevented or treated by the pharmaceutical compositions of this invention include cognitive pathologies such as dementia, cerebral ischemia, trauma, epilepsy, schizophrenia, chronic pain and neurologic disorders such as but not limited to depression, social phobia and obsessive compulsive disorders.

Cardiovascular disorders to be prevented or treated by the pharmaceutical compositions of this invention include ischemic disorders, infarct or reperfusion damage, atherosclerosis and stroke.

Allergic conditions to be prevented or treated by the pharmaceutical compositions of this invention include those caused by the pollen of gramineae, the presence of pets, as well as more severe forms, such as asthma, characterized by inflammation of airways and bronchospasm. Without wishing to be bound by theory, the antiallergic effect of the compounds of the invention may be related to their suppression of certain B-cell activation pathways, which can lead to the suppression of IgE release. It may also be related to their properties of inhibiting certain Th2 cytokines, such as IL-5, IL-13 or IL-10, involved in asthma.

TNF- α -related disorders to be prevented or treated by the pharmaceutical compositions of this invention include the following:

- septic or endotoxic shock or sepsis, especially in patients with a serum level of interleukin-6 above 1,000 pg/ml at start of treatment;
- vascular TNF- α - mediated diseases such as, but not limited to, disseminated intravascular coagulation and Kawasaki's pathology;
- pathologies and conditions associated with and/or induced by abnormal levels of TNF- α (herein defined as exceeding by at least 10 % and at most 500% the TNF- α level present in a normal healthy subject) occurring in a systemic, localized or particular tissue type or location in the body of the mammal; such tissue types include, but are not limited to, blood, lymph, liver, kidney, spleen, heart muscle or blood vessels, brain or spinal cord white matter or grey matter, cartilage, ligaments, tendons, lung, pancreas, ovary, testes and prostate. Abnormal TNF levels can also be localized to specific regions or cells in the body, such as joints, nerve blood vessel junctions and bones. Such pathologies include alcohol-induced hepatitis; neurodegenerative diseases such as extrapyramidal and cerebellar disorders including lesions of the corticospinal system; disorders of the basal ganglia; hyperkinetic movement disorders such as chorea; drug-induced movement disorders; hypokinetic movement disorders,

- such as Parkinson's disease; spinocerebellar degenerations such as spinal ataxia, multiple systems degenerations (including Dejerine-Klumpke syndrome) and systemic disorders (including Refsum's disease, abetalipoproteinemia, ataxia and telangiectasia); disorders of the motor unit, such as neurogenic muscular atrophies (anterior horn cell degeneration, such as amyotrophic lateral sclerosis, infantile spinal muscular atrophy and juvenile spinal muscular atrophy); Alzheimer's disease; Wernicke-Korsakoff syndrome; Creutzfeldt-Jakob disease; Hallerorden-Spatz disease; and primary or secondary myelodysplastic syndromes;
- toxic effects of TNF- α and/or anti-cancer chemotherapeutic agents, especially side effects associated with TNF generation during neoplastic therapy, for instance following use of cisplatin;
 - injuries after irradiation of a tissue of a mammal by radio-elements, such as but not limited to radiation-induced graft-versus-host disease; and
 - cachexia and similar chronic wasting diseases, whether associated with cancer or with other chronic diseases such as malabsorptive disorders, excessive physical stress, eating disorders and AIDS.

The medicament of this invention may be for prophylactic use, i.e. where circumstances are such that an elevation in the TNF level might be expected or alternatively, may be for use in reducing the TNF level after it has reached an undesirably high level or as the TNF level is rising.

The term "pharmaceutically acceptable carrier or excipient" as used herein in relation to pharmaceutical compositions and combined preparations means any material or substance with which the active principle, i.e. the pteridine derivative of the general formula (I) or the general formula (V), and optionally the immunosuppressant or immunomodulator or antineoplastic drug or antiviral agent, may be formulated in order to facilitate its application or dissemination to the locus to be treated, for instance by dissolving, dispersing or diffusing the said composition, and/or to facilitate its storage, transport or handling without impairing its effectiveness. The pharmaceutically acceptable carrier may be a solid or a liquid or a gas which has been compressed to form a liquid, i.e. the compositions of this invention can suitably be used as concentrates, emulsions, solutions, granulates, dusts, sprays, aerosols, pellets or powders.

Suitable pharmaceutical carriers for use in the said pharmaceutical compositions and their formulation are well known to those skilled in the art. There is no particular restriction to their selection within the present invention although, due to the usually low or very low water-solubility of the pteridine derivatives of this invention, special attention will be paid to the selection of suitable carrier combinations that can assist in properly formulating them in view of the expected time release profile. Suitable pharmaceutical carriers include additives such as wetting agents, dispersing agents, stickers, adhesives, emulsifying or surface-active agents, thickening agents, complexing agents, gelling agents, solvents, coatings, antibacterial

and antifungal agents (for example phenol, sorbic acid, chlorobutanol), isotonic agents (such as sugars or sodium chloride) and the like, provided the same are consistent with pharmaceutical practice, i.e. carriers and additives which do not create permanent damage to mammals. The pharmaceutical compositions of the present invention may be prepared in any
5 known manner, for instance by homogeneously mixing, dissolving, spray-drying, coating and/or grinding the active ingredients, in a one-step or a multi-steps procedure, with the selected carrier material and, where appropriate, the other additives such as surface-active agents. may also be prepared by micronisation, for instance in view to obtain them in the form of microspheres usually having a diameter of about 1 to 10 μm , namely for the manufacture of
10 microcapsules for controlled or sustained release of the biologically active ingredient(s).

Suitable surface-active agents to be used in the pharmaceutical compositions of the present invention are non-ionic, cationic and/or anionic materials having good emulsifying, dispersing and/or wetting properties. Suitable anionic surfactants include both water-soluble soaps and water-soluble synthetic surface-active agents. Suitable soaps are alkaline or
15 alkaline-earth metal salts, unsubstituted or substituted ammonium salts of higher fatty acids ($\text{C}_{10}\text{-C}_{22}$), e.g. the sodium or potassium salts of oleic or stearic acid; or of natural fatty acid mixtures obtainable from coconut oil or tallow oil. Synthetic surfactants include sodium or calcium salts of polyacrylic acids; fatty sulphonates and sulphates; sulphonated benzimidazole derivatives and alkylarylsulphonates. Fatty sulphonates or sulphates are
20 usually in the form of alkaline or alkaline-earth metal salts, unsubstituted ammonium salts or ammonium salts substituted with an alkyl or acyl radical having from 8 to 22 carbon atoms, e.g. the sodium or calcium salt of lignosulphonic acid or dodecylsulphonic acid or a mixture of fatty alcohol sulphates obtained from natural fatty acids, alkaline or alkaline-earth metal salts of sulphuric or sulphonic acid esters (such as sodium lauryl sulphate) and sulphonic acids of
25 fatty alcohol/ethylene oxide adducts. Suitable sulphonated benzimidazole derivatives preferably contain 8 to 22 carbon atoms. Examples of alkylarylsulphonates are the sodium, calcium or alcanolamine salts of dodecylbenzene sulphonic acid or dibutyl-naphtalenesulphonic acid or a naphtalene-sulphonic acid/formaldehyde condensation product. Also suitable are the corresponding phosphates, e.g. salts of phosphoric acid ester and an
30 adduct of p-nonylphenol with ethylene and/or propylene oxide, or phospholipids. Suitable phospholipids for this purpose are the natural (originating from animal or plant cells) or synthetic phospholipids of the cephalin or lecithin type such as e.g. phosphatidylethanolamine, phosphatidylserine, phosphatidylglycerine, lysolecithin, cardiolipin, dioctanyl-phosphatidylcholine, dipalmitoylphosphatidylcholine and their mixtures.

35 Suitable non-ionic surfactants include polyethoxylated and polypropoxylated derivatives of alkylphenols, fatty alcohols, fatty acids, aliphatic amines or amides containing at least 12 carbon atoms in the molecule, alkylarenesulphonates and dialkylsulphosuccinates, such as polyglycol ether derivatives of aliphatic and cycloaliphatic alcohols, saturated and unsaturated fatty acids and alkylphenols, said derivatives preferably containing 3 to 10 glycol

ether groups and 8 to 20 carbon atoms in the (aliphatic) hydrocarbon moiety and 6 to 18 carbon atoms in the alkyl moiety of the alkylphenol. Further suitable non-ionic surfactants are water-soluble adducts of polyethylene oxide with polypropylene glycol, ethylenediaminopolypropylene glycol containing 1 to 10 carbon atoms in the alkyl chain, which
5 adducts contain 20 to 250 ethyleneglycol ether groups and/or 10 to 100 propyleneglycol ether groups. Such compounds usually contain from 1 to 5 ethyleneglycol units per propyleneglycol unit. Representative examples of non-ionic surfactants are nonylphenol-polyethoxyethanol, castor oil polyglycolic ethers, polypropylene/ polyethylene oxide adducts, tributylphenoxypolyethoxyethanol, polyethyleneglycol and octylphenoxypolyethoxyethanol.
10 Fatty acid esters of polyethylene sorbitan (such as polyoxyethylene sorbitan trioleate), glycerol, sorbitan, sucrose and pentaerythritol are also suitable non-ionic surfactants.

Suitable cationic surfactants include quaternary ammonium salts, preferably halides, having 4 hydrocarbon radicals optionally substituted with halo, phenyl, substituted phenyl or hydroxy; for instance quaternary ammonium salts containing as N-substituent at least one C₈-
15 C₂₂ alkyl radical (e.g. cetyl, lauryl, palmityl, myristyl, oleyl and the like) and, as further substituents, unsubstituted or halogenated lower alkyl, benzyl and/or hydroxy-lower alkyl radicals.

A more detailed description of surface-active agents suitable for this purpose may be found for instance in " McCutcheon's Detergents and Emulsifiers Annual " (MC Publishing Corp., Ridgewood, New Jersey, 1981), " Tensid-Taschenbuch ", 2nd ed. (Hanser Verlag,
20 Vienna, 1981) and "Encyclopaedia of Surfactants (Chemical Publishing Co., New York, 1981).

Structure-forming, thickening or gel-forming agents may be included into the pharmaceutical compositions and combined preparations of the invention. Suitable such agents are in particular highly dispersed silicic acid, such as the product commercially
25 available under the trade name Aerosil; bentonites; tetraalkyl ammonium salts of montmorillonites (e.g., products commercially available under the trade name Bentone), wherein each of the alkyl groups may contain from 1 to 20 carbon atoms; cetostearyl alcohol and modified castor oil products (e.g. the product commercially available under the trade name Antissettle).

30 Gelling agents which may be included into the pharmaceutical compositions and combined preparations of the present invention include, but are not limited to, cellulose derivatives such as carboxymethylcellulose, cellulose acetate and the like; natural gums such as arabic gum, xanthum gum, tragacanth gum, guar gum and the like; gelatin; silicon dioxide; synthetic polymers such as carbomers, and mixtures thereof. Gelatin and modified celluloses
35 represent a preferred class of gelling agents.

Other optional excipients which may be included in the pharmaceutical compositions and combined preparations of the present invention include additives such as magnesium oxide; azo dyes; organic and inorganic pigments such as titanium dioxide; UV-absorbers; stabilisers;

odor masking agents; viscosity enhancers; antioxidants such as, for example, ascorbyl palmitate, sodium bisulfite, sodium metabisulfite and the like, and mixtures thereof; preservatives such as, for example, potassium sorbate, sodium benzoate, sorbic acid, propyl gallate, benzylalcohol, methyl paraben, propyl paraben and the like; sequestering agents such as ethylene-diamine tetraacetic acid; flavoring agents such as natural vanillin; buffers such as citric acid and acetic acid; extenders or bulking agents such as silicates, diatomaceous earth, magnesium oxide or aluminum oxide; densification agents such as magnesium salts; and mixtures thereof.

Additional ingredients may be included in order to control the duration of action of the biologically-active ingredient in the compositions and combined preparations of the invention. Control release compositions may thus be achieved by selecting appropriate polymer carriers such as for example polyesters, polyamino-acids, polyvinyl-pyrrolidone, ethylene-vinyl acetate copolymers, methylcellulose, carboxymethylcellulose, protamine sulfate and the like. The rate of drug release and duration of action may also be controlled by incorporating the active ingredient into particles, e.g. microcapsules, of a polymeric substance such as hydrogels, polylactic acid, hydroxymethyl-cellulose, polymethyl methacrylate and the other above-described polymers. Such methods include colloid drug delivery systems like liposomes, microspheres, microemulsions, nanoparticles, nanocapsules and so on. Depending on the route of administration, the pharmaceutical composition or combined preparation of the invention may also require protective coatings.

Pharmaceutical forms suitable for injectable use include sterile aqueous solutions or dispersions and sterile powders for the extemporaneous preparation thereof. Typical carriers for this purpose therefore include biocompatible aqueous buffers, ethanol, glycerol, propylene glycol, polyethylene glycol, complexing agents such as cyclodextrins and the like, and mixtures thereof.

Since, in the case of combined preparations including the pteridine derivative of this invention and an immunosuppressant or immunomodulator or antihistamine or antineoplastic drug or antiviral agent, both ingredients do not necessarily bring out their synergistic therapeutic effect directly at the same time in the patient to be treated, the said combined preparation may be in the form of a medical kit or package containing the two ingredients in separate but adjacent form. In the latter context, each ingredient may therefore be formulated in a way suitable for an administration route different from that of the other ingredient, e.g. one of them may be in the form of an oral or parenteral formulation whereas the other is in the form of an ampoule for intravenous injection or an aerosol.

The present invention further relates to a method for preventing or treating a disease selected from the group consisting of CNS disorders, cell proliferative disorders, allergic conditions, viral infections, immune and auto-immune disorders, transplant rejections and TNF- α -related disorders in a patient, preferably a mammal, more preferably a human being. The method of this invention consists of administering to the patient in need thereof an

- effective amount of a pteridine derivative having the general formula (I) or the general formula (V), optionally together with an effective amount of another immunosuppressant or immunomodulator or antineoplastic drug or antiviral agent, or a pharmaceutical composition comprising the same, such as disclosed above in extensive details. The effective amount is usually in the range of 0.01 mg to 20 mg, preferably 0.1 mg to 5 mg, per day per kg bodyweight for humans. Depending upon the pathologic condition to be treated and the patient's condition, the said effective amount may be divided into several sub-units per day or may be administered at more than one day intervals. The patient to be treated may be any warm-blooded animal, preferably a human being, suffering from said pathologic condition.
- The following examples are intended to illustrate several embodiments of the present invention, including the preparation of the pteridine derivatives and their pyrimidine intermediates, without limiting its scope in any way.

Example 1 – preparation of 2,6-diamino-5-nitroso-4-hydroxypyrimidine

- The following illustrates the method step (a) shown in figure 1. To a solution of 2,6-diamino-4-hydroxypyrimidine (12.9 g, 102 mmoles) in 200 ml of a 10% acetic acid solution in water at 80 °C was added dropwise a solution of NaNO₂ (7.05 g, 102 mmoles) in 20 ml water. A pink precipitate was formed, which was further stirred for 1 hour at 80 °C. The reaction mixture was cooled down in the refrigerator overnight. The precipitate was filtered off and dried over P₂O₅, yielding the title compound as a pink powder (15.43 g, 97%). Its spectral data are in accordance with literature data (Traube in *Ber.* (1900) 33:1371 and Landauer et al. in *J. Chem. Soc.* (1953) 3721-3722).

Example 2 - synthesis of 2,5,6-triamino-4-hydroxypyrimidine

- The following illustrates the method step (b) shown in figure 1. A suspension of the compound of example 1 (15 g, 96.7 mmoles) in an ammonium sulfide solution (20 % in water, 200 ml) was stirred overnight at 50 °C. The reaction mixture was cooled down in the refrigerator and the formed precipitate was filtered off, yielding the title compound as a yellow powder (11.33 g, 83 %). The spectral data are identical with literature data (same as for example 1).

Example 3 - synthesis of 3,4-dimethoxyphenylglyoxalmonoxime

- In a mixture of dioxane (250 ml) and water (10 ml), SeO₂ (0.33 mole) was heated to 50 °C. After solution of SeO₂, 3,4-dimethoxyacetophenone was added and the mixture heated under reflux for 16 hours. The hot solution was filtered to remove selenium. The filtrate was evaporated, the oily residue dissolved in CHCl₃ (300 ml), then washed with saturated NaHCO₃ solution (100 ml) and water. The organic phase was dried over Na₂S₂O₄, filtered and evaporated. The yellow oil was distilled in vacuum, the resulting 3,4-dimethoxyphenylglyoxal was dissolved in MeOH (50 ml) and water (200 ml), then acetoxime (0.25 mol) was added

and the pH adjusted to 4 by 2 N HCl. The solution was heated to 50 °C for 2 hours, then cooled to 0 °C and the resulting crystals collected. After washing with cold water and drying in a vacuum desiccator, 3,4-dimethoxyphenylglyoxalmonooxime was obtained with a yield of 71 %. Recrystallization can be achieved from CHCl₃ or acetone. The compound was further
5 characterized by nuclear magnetic resonance spectra as follows: ¹H-NMR (200 MHz, DMSO-d₆): δ 3.80 (3 H, s), 3.84 (3 H, s), 7.06 (1 H, d), 7.51 (1 H, s), 7.75 (1 H, d), 8.10 (1 H, s) and 12.51 (1 H, s) ppm.

Example 4 - synthesis of 2-amino-6-(3,4-dimethoxyphenyl) pterine

10 The following illustrates the method step (c) shown in figure 1. To a boiling solution of the compound of example 2 (2.4 g, 17 mmole) in methanol (100 ml, with 0.9 N HCl) was added dropwise a solution of the compound of example 3 (3.8 g, 18.2 mmole) in methanol (100 ml). The reaction mixture was heated under reflux for 4 hours. A precipitate was formed, which was filtered off, washed with water, ethanol and diethyl ether. The precipitate was dried over
15 P₂O₅ under vacuum, yielding the title compound as a yellow powder (4.33 g, 85 %). This compound was further characterized by nuclear magnetic resonance spectra as follows:

- ¹H-NMR (500 MHz, TFA): δ 4.11 (3 H, s), 4.07 (3 H, s), 7.21 (1 H, d), 7.78 (1 H, dd), 7.81 (1 H, d) and 9.32 (1 H, s) ppm.
- ¹³C-NMR (125 MHz, TFA): δ at 56.39, 56.76, 111.94, 113.21, 123.22, 127.41, 127.91,
20 145.92, 149.39, 150.46, 152.47, 153.15, 155.13 and 161.59 ppm.

Example 5 - synthesis of 2-acetylamino-6-(3,4-dimethoxyphenyl) pterine

A suspension of the compound of example 4 (10.46 g, 35 mmole) in acetic anhydride (600 ml) and acetic acid (200 ml) was refluxed for 1 hour until a clear solution was formed. By
25 cooling down the reaction mixture in a refrigerator, a precipitate was formed which was filtered off, washed with ethyl acetate and diethyl ether. The precipitate was dried over P₂O₅ under vacuum, yielding the title compound as a yellow powder (9.19 g, 77 %). This compound was further characterized as follows:

- mass spectrum (MS): m/z (%): 300 ([M+H]⁺, 100)
- 30 - ¹H-NMR (200 MHz, DMSO-d₆): δ 2.22 (3 H, s), 3.84 (3 H, s), 3.87 (3 H, s), 7.14 (1 H, d), 7.75 (2 H, m) and 9.51 (1 H, s) ppm.

Example 6 - synthesis of 2-acetylamino-4-(1,2,4-triazolyl)-6-[3,4-(dimethoxy-phenyl)] pteridine

35 The following illustrates the method step (f) shown in figure 1. To a solution of phosphorus oxychloride (1.68 ml, 18 mmole) and 1,2,4-triazole (4.96 g, 72 mmole) in dry pyridine (110 ml) was added the compound of example 5 (2.45 g, 7.2 mmole). The suspension was stirred at room temperature for 4 hours. The precipitate was filtered off, washed with pyridine, toluene and diethyl ether. The resulting solid was dried over P₂O₅ under

vacuum, affording the title compound as a yellow powder (2 g, yield: 80 %) characterized by mass spectrum data as follows: m/z (%): 392 ($[M+H]^+$, 100).

Example 7 - synthesis of 2-acetylamino-4-(piperazin-1-yl)-6-(3,4-dimethoxy-phenyl) pteridine

5 The following illustrates the method step (g) shown in figure 1. To a suspension of the compound of example 6 (3.92 g, 10 mmole) in dioxane (200 ml) was added piperazine (1.29 g, 15 mmole). The suspension was stirred for 16 hours at room temperature. The precipitate was filtered off and washed with dioxane, ethanol and diethyl ether. The solid was dried over P_2O_5 under vacuum, yielding the title compound as a yellow powder (3.48 g, 85 %).

10

Examples 8 to 21 - synthesis of 2-amino-4-(N-acyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridines

The following illustrates the method step (h) shown in figure 1. To a suspension of the compound of example 7 (409 mg, 1 mmole) in dry pyridine (40 ml) was added under nitrogen
15 a suitable carboxylic acid chloride (1.2 mmole). The suspension was stirred at room temperature for 24 hours. The solvent was concentrated *in vacuo* and the crude residue was dissolved in a mixture of $CH_3OH/20\%$ K_2CO_3 in water (1:1). The solution was stirred at room temperature for 16 hours. Evaporation of the solvents *in vacuo*, followed by purification of the residue by preparative TLC (silica, using a CH_3OH/CH_2Cl_2 mixture (5:95) as the eluent)
20 afforded the desired compound as a yellow powder. This procedure provided, with a yield ranging from 33 % to 75 % depending upon the carboxylic acid chloride used, the following pure final pteridine derivatives which were characterized by their mass spectrum MS and optionally by their 1H -NMR (500 MHz, $DMSO-d_6$) spectrum:

- 2-amino-4-(N-acetylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (example 8): MS
25 410 ($[M+H]^+$;
- 2-amino-4-[(N-propionyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 9): MS 424 ($[M+H]^+$;
- 2-amino-4-[(N-hexanoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 10): MS 466 ($[M+H]^+$;
- 30 - 2-amino-4-(N-benzoylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 11): MS 472 ($[M+H]^+$; 1H -NMR: δ at 3.65 (br s, 2 H), 3.80-3.90 (m, 8 H), 4.37 (br s, 4 H), 6.76 (br s, 2 H, NH_2), 7.07 (d, 1 H), 7.48 (m, 5 H), 7.59 (br d, 1 H), 7.66 (dd, 1 H) and 9.31 (s, 1 H) ppm;
- 2-amino-4-[N-(4-chlorobenzoyl)]piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine
35 (example 12): MS 506 ($[M+H]^+$;
- 2-amino-4-[(N-2-thiophenecarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 13): MS 478 ($[M+H]^+$;
- 2-amino-4-[(N-diethylcarbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 14): MS 467 ($[M+H]^+$;

- 2-amino-4-[(*N*-hydrocinnamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 15): MS 500 ($[M+H]^+$);
- 2-amino-4-[(*N*-(4-cyanobenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 16): MS 497 ($[M+H]^+$);
- 5 - 2-amino-4-[(*N*-phenoxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 17): MS 502 ($[M+H]^+$; 1H -NMR δ at 3.75 (br d, 4 H), 3.83 (s, 3 H), 3.86 (s, 3 H), 4.32 (br s, 2 H), 4.41 (br s, 2 H), 4.90 (s, 2H), 6.77 (br s, 2 H), 6.94-6.97 (m, 3 H), 7.09 (d, 1 H), 7.28-7.31 (m, 2 H), 7.62 (d, 1 H), 7.67 (dd, 1 H) and 9.32 (s, 1 H) ppm ;
- 10 - 2-amino-4-[(*N*-4-butylbenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 18): MS 528 ($[M+H]^+$);
- 2-amino-4-[(*N*-isonicotinoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 19): MS 473 ($[M+H]^+$; 1H -NMR δ at 3.58 (br s, 2 H), 3.82 (s, 6 H), 3.87 (br s, 2 H), 4.33 (br s, 2 H), 4.40 (br s, 2 H), 6.77 (br s, 2 H), 7.07 (d, 1 H), 7.47 (dd, 2 H), 7.59 (br d), 7.66 (dd, 1 H), 8.71 (dd, 2 H) and 9.32 (s, 1 H) ppm;
- 15 - 2-amino-4-[(*N*-diisopropylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 20): MS 495 ($[M+H]^+$; and
- 2-amino-4-[(*N*-(4-pentoxycarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 21): MS 558 ($[M+H]^+$).

20 Example 22 - synthesis of 2,6-diamino-4-chloro-5-*p*-chlorophenylazopyrimidine

The following illustrates the method step (a) shown in figure 2. A solution of *p*-chloroaniline (25.5 g, 0.2 mole) in 6 N HCl (100 ml) was cooled to 0 °C and then NaNO₂ (13.8 g, 0.2 mole) in water (40 ml) was added dropwise with stirring. After the addition was completed, the solution was stirred for another 30 minutes. Urea (5 g) was added to destroy

25 the excess of HNO₂. The diazonium salt solution was then poured into a solution of 2,6-diamino-4-chloropyrimidine (26 g, 0.18 mole) in water (500 ml) and stirred for 30 minutes. Then potassium acetate (70 g) was added and the mixture was stirred for 16 hours at room temperature. The resulting precipitate was collected by suction, washed with water and dried in a vacuum dessicator over P₂O₅ to give 44 g (81 %) of a yellow solid. Recrystallization can

30 be achieved from DMF/H₂O. The spectral data are in analogy with those described in literature (Fröhlich et al. in *J. Med. Chem.* (1999) 42:4108-4121).

Example 23 - synthesis of 2,6-diamino-4-(piperazin-1-yl)-5-*p*-chlorophenylazo-pyrimidine

The following illustrates the method step (b) shown in figure 2. A solution of the

35 compound of example 22 (5 g, 16.6 mole) in DMF (50 ml) and piperazine (10 g) was heated in an oilbath to 70 °C for 5 hours. Water (50 ml) was added and the reaction mixture was cooled down. The yellow precipitate was filtered off, washed with water and dried. Recrystallization can be achieved from ethanol.

Example 24 - synthesis of 2,5,6-triamino-4-(piperazin-1-yl) pyrimidine

The following illustrates the method step (c) shown in figure 2. To a suspension of the compound of example 23 (2.03 g, 6.1 mmole) in ethanol (98 ml) and water (98 ml) was added zinc (4.04 g) and acetic acid (2.02 ml). The suspension was refluxed until a clear solution was obtained, i.e. for 1 hour. Zinc was filtered off and the filtrate was evaporated, followed by co-evaporation with toluene. A brown precipitate was obtained, which was resuspended in diethyl ether and stirred overnight at room temperature in order to remove *p*-chloroaniline. The precipitate was filtered off and dried in a vacuum dessicator over P₂O₅.

Example 25 - synthesis of 2-amino-4-(piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine

The following illustrates the method step (d) shown in figure 2. To a solution of the compound of example 24 (2.65 mmole, 554 mg) in methanol (30 ml) was added 3,4-dimethoxyphenylglyoxaloxime (2.65 mmole, 554 mg). The pH of the reaction mixture was adjusted to 3 by adding a few drops of concentrated hydrochloric acid. The mixture was refluxed for 3 hours. A yellow precipitate was formed. The reaction mixture was cooled down and neutralized by the addition of concentrated NH₃ till pH 9. The precipitate was filtered off and used for further reaction without any purification.

Examples 26 to 32 - synthesis of 2-amino-4-(*N*-acylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridines and 2-amino-4-(*N*-sulfonylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridines

The following illustrates the method step (g) shown in figure 2. To a suspension of the crude compound of example 25 (506 mg, 1.38 mmole) in pyridine (30 ml) was added a suitable carboxylic acid chloride or sulfonyl chloride (2.07 mmole). The suspension was stirred at room temperature overnight. After pyridine evaporation, the residue was purified by flash chromatography on silica gel, the mobile phase being CH₃OH/CH₂Cl₂ mixtures (in a ratio gradually ranging from 2:98 to 5:95), optionally followed by preparative TLC on silica gel (the mobile phase being a 7:93 CH₃OH/CH₂Cl₂ mixture), thus affording the desired compound as a yellow powder. This procedure provided, with a yield ranging from 30 % to 50 % depending upon the carboxylic acid or sulfonyl chloride used, the following pure final pteridine derivatives which were characterized by their mass spectrum MS and optionally by their ¹H-NMR (200 MHz, DMSO-*d*₆) spectrum:

- 2-amino-4-[*N*-(3-methoxybenzoyl)piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 26): MS 502 ([M+H]⁺; ¹H-NMR: 3.33 (br s, 4 H), 3.80 (s, 9 H), 4.37 (br s, 4 H), 6.78 (br s, 2 H), 6.99-7.70 (m, 7 H) and 9.33 (s, 1 H) ppm.
- 2-amino-4-[*N*-(2-furoyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 27): MS 462 ([M+H]⁺;
- 2-amino-4-[*N*-(benzyloxyacetyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 28): MS 516 ([M+H]⁺;

- 2-amino-4-[(N-(p-chlorophenoxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 29): MS 536 ([M+H]⁺;
- 2-amino-4-[(N-cyclohexylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 30): MS 478 ([M+H]⁺;
- 5 - 2-amino-4-[(N-phenylsulfonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 31): MS 508 ([M+H]⁺; ¹H-NMR: 3.14 (br s, 4 H), 3.84 (s, 3 H), 3.87 (s, 3 H), 4.40 (br s, 4 H), 6.88 (br s, 2 H), 7.09 (d, 1 H) and 7.56-7.78 (m, 7 H) ppm ; and
- 2-amino-4-[(N-p-fluorobenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 32): MS 490 ([M+H]⁺.

10

Example 33 - synthesis of 2,5,6-triamino-4-hydroxypyrimidine dihydrochloride

The following illustrates the method step (a) shown in figure 3. To a stirred suspension of 2,4,5-triamino-6-hydroxy-pyrimidine (40.93 g, 290 mmole) in methanol (500 ml) was added dropwise concentrated hydrochloric acid 37% (60.5 ml, 725 mmole). The suspension was stirred 30 minutes at room temperature, filtered and the precipitate was washed with methanol, diethyl ether and dried over KOH under vacuum, affording the title compound as a white powder (53.82 g, yield 88 %). Spectral data are identical with literature data (W. Pfeiderer, *Chem. Ber.* (1957) 90:2272).

20

Example 34 - synthesis of 2-amino-6-(3,4-dimethoxyphenyl)pterine

The following illustrates the method step (b) shown in figure 3. The compound of example 33 (21.5 g, 102 mmole) and 3,4-dimethoxyphenyl-glyoxalmonooxime (25.61 g, 122.4 mmole) were suspended in methanol (400 ml) and the orange suspension was heated under reflux for 2.5 hours. The yellow suspension was cooled with an ice bath and the precipitate was filtered and successively washed with methanol, ethyl acetate, diethyl ether, and dried at 110°C for 3 hours to afford a shiny yellow powder (21.56 g, yield 71 %) which was used without further purification.

25

Example 35 - preparation of 2-amino-4-(piperazin-1-yl)-6-(3,4-dimethoxy-phenyl) pteridine

30

The following illustrates the method step (e) shown in figure 3. Piperazine (12.06 g, 140 mmole) was added to a stirred suspension of the compound of example 34 (5.99 g, 20 mmole) in pyridine (64 ml) and 1,1,1,3,3,3-hexamethyldisilazane (64 ml, 300 mmole) in the presence of a catalytic amount of ammonium sulfate and *p*-toluenesulfonic acid. The mixture was heated under reflux for 15 hours and the brown solution was cooled with an ice bath. Methanol was added and the mixture was evaporated to dryness. The brown residue was co-evaporated 2 times with xylene, and adsorbed on silica gel. The compound was purified on silica gel column chromatography, using a 9/1 CH₂Cl₂/MeOH mixture containing 1 % concentrated aqueous ammonia as eluent, thus affording the desired compound (3.12 g, yield

35

42 %) which was characterized by its mass spectrum MS and $^1\text{H-NMR}$ (200 MHz, $\text{DMSO-}d_6$) spectrum as follows:

- MS: m/z (%): 368 ($[\text{M}+\text{H}]^+$, 100); and
- $^1\text{H-NMR}$ (200 MHz, $\text{DMSO-}d_6$): δ 2.92 (4 H, br s), 3.83 (3 H, s), 3.86 (3 H, s), 4.27 (4 H, br s), 6.70 (2 H, br s), 7.08 (1 H, d), 7.62-7.69 (2 H, m) and 9.30 (1 H, s) ppm.

Examples 36 to 51 - synthesis of 2-amino-4-(*N*-acylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridines, 2-amino-4-(*N*-sulfonylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridines and 2-amino-4-(*N*-thioacylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridines

The following illustrates the method step (f) shown in figure 3. To a suspension of the compound of example 35 (184 mg, 0.5 mmole) in CH_2Cl_2 (10 ml) was added triethylamine (84 μl , 0.6 mmole) and a suitable carboxylic acid, thiocarboxylic or sulfonyl chloride (0.55 mmole). The suspension was stirred at room temperature for 2 to 24 hours. The solution was diluted CH_2Cl_2 (20 ml) and extracted with a 5% aqueous solution of sodium hydrogen carbonate (30 ml). The organic layer was concentrated *in vacuo* and the crude residue was subjected to silica gel column chromatography, the mobile phase being $\text{CH}_3\text{OH}/\text{CH}_2\text{Cl}_2$ mixtures (in a ratio gradually ranging from 1:99 to 5:95), thus affording the desired compound as a yellow powder. This procedure provided, with a yield ranging from 35 % to 85 % depending upon the carboxylic acid, thiocarboxylic or sulfonyl chloride used, the following pure final pteridine derivatives which were characterized by their mass spectrum MS:

- 2-amino-4-[(*N*-2-thiophenacetyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 36): MS 492 ($[\text{M}+\text{H}]^+$;
- 2-amino-4-[(*N*-cinnamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 37): MS 498 ($[\text{M}+\text{H}]^+$;
- 2-amino-4-[(*N*-1-pyrrolidinylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 38): MS 951 ($[\text{2M}+\text{Na}]^+$, 15); 929 ($[\text{2M}+\text{H}]^+$, 15); 465 ($[\text{M}+\text{H}]^+$, 100)
- 2-amino-4-[(*N*-diphenylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 39): MS 563 ($[\text{M}+\text{H}]^+$;
- 2-amino-4-[*N*-(2,6-dichloro-5-fluoro-nicotinoyl)]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 40): MS 559 ($[\text{M}+\text{H}]^+$;
- 2-amino-4-[(*N*-methoxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 41): MS 901 ($[\text{2M}+\text{Na}]^+$, 20); 879 ($[\text{2M}+\text{H}]^+$, 10) and 440 ($[\text{M}+\text{H}]^+$, 100);
- 2-amino-4-[*N*-(2-methoxybenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 42): MS 502 ($[\text{M}+\text{H}]^+$;
- 2-amino-4-[(*N*-benzylsulfonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 43): MS 522 ($[\text{M}+\text{H}]^+$;
- 2-amino-4-[*N*-(3,4-dichlorobenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 44): MS 540 ($[\text{M}+\text{H}]^+$;

- 2-amino-4-[*N*-(4-chlorophenylacetyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 45): MS 520 ($[M+H]^+$;
- 2-amino-4-[*N*-(1-naphthoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 46): MS 522 ($[M+H]^+$;
- 5 - 2-amino-4-[*N*-(3-furoylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 47): MS 490 ($[M+H]^+$;
- 2-amino-4-[*N*-(benzyloxycarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 48): MS 502 ($[M+H]^+$;
- 2-amino-4-[*N*-(dimethylthiocarbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 49): MS 455 ($[M+H]^+$;
- 10 - 2-amino-4-[*N*-(phenoxy carbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 50): MS 487 ($[M+H]^+$; and
- 2-amino-4-[*N*-(phenoxythiocarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 51): MS 504 ($[M+H]^+$.

15

Example 52 - synthesis of 2,6-diamino-4-(*N*-acetyl piperazin-1-yl)-pyrimidine

The following illustrates the method step (a) shown in figure 4. *N*-acetyl piperazine (12.82 g, 100 mmole) was added to a stirred suspension of 4-chloro-2,6-diaminopyrimidine (7.23 g, 50 mmole, m.p. 199°C, commercially available for instance from Merck or from Qiaoj

20 Group Co. Ltd., Hong-Kong) in water (100 ml), and the mixture was refluxed for 21 hours. The orange solution was cooled and made alkaline with NaOH 10M (5 ml) to lead to a white precipitation. The solution was filtered; the solid was washed with cold water and dried over P_2O_5 in a vacuum dessicator to afford the desired compound as a white powder (9.77 g, yield 82%) which was characterized by the following mass spectrum MS m/z (%): 237 ($[M+H]^+$, 100); 195 ($[M-Ac+H]^+$, 25).

25

Example 53 - synthesis of 2,6-diamino-5-nitroso-4-(*N*-acetyl piperazin-1-yl)-pyrimidine

The following illustrates the method step (b) shown in figure 4. Acetic acid (4 ml) was added dropwise to a stirred suspension of the compound of example 52 (9.45 g, 40 mmole)

30 and sodium nitrite (3.04 g, 44 mmole) in water (200 ml) at room temperature. The purple mixture was stirred for 1 hour and cooled down to 5°C for 14 hours. The precipitate was filtered, washed with water, diethyl ether and dried over P_2O_5 in a vacuum dessicator to afford the titled compound as a purple powder (10.53 g, yield 99%) which was characterized by the following mass spectrum MS m/z (%): 288 ($[M+Na]^+$, 60); 266 ($[M+H]^+$, 100).

35

Example 54 - synthesis of 2,5,6-triamino-4-(*N*-acetyl piperazin-1-yl)-pyrimidine

The following illustrates the method step (c) shown in figure 4. To a suspension of the compound obtained in example 53 (1 g, 3.77 mmole) in water (25 ml) was added sodium dithionite (1.97 g, 11.3 mmole). The suspension was heated to 50 °C till a clear solution was

obtained. Water was evaporated *in vacuo* and the residue was co-evaporated with toluene twice. The crude material was used for further reaction without any purification.

Example 55 - synthesis of 2-amino-4-(N-acetylpiperazin-1-yl)-6-phenyl-pteridine

5 The following illustrates the method step (d) shown in figure 4. The crude product obtained in example 54 and isonitrosoacetophenone (653 mg, 4.0 mmole) were suspended in a 1.25 M HCl solution in MeOH (20 ml) and the mixture was refluxed for 3 hours. The reaction mixture was cooled down to room temperature and neutralized with a 25 % aqueous NH₃ solution to pH 9. The mixture was evaporated to dryness and the residue was partitioned
10 between CHCl₃ and H₂O. The organic layer was separated, evaporated to dryness and purified by silica gel chromatography, the mobile phase consisting of CH₃OH/CH₂Cl₂ mixtures (in a ratio gradually ranging from 2:98 to 4:96), thus affording the desired compound as a yellow powder (978 mg, yield 70 %) was characterized by its mass spectrum MS as follows: MS *m/z* (%): 721 ([2M+Na]⁺, 60); 372 ([M+Na]⁺, 10) and 350 ([M+H]⁺, 100).

Example 56 - synthesis of 2-amino-4-(N-acetylpiperazin-1-yl)-6-(4-tolyl) pteridine

15 The method of example 55 was repeated, except for using 4-methylphenylglyoxalmonoxime (4.0 mmole) instead of isonitrosoaceto-phenone. This afforded the title compound as a yellow powder (900 mg, yield 62 %), which was characterized by its
20 mass spectrum MS as follows: MS *m/z* (%): 362 ([M+H]⁺, 100).

Example 57 - synthesis of 2-amino-4-(N-acetylpiperazin-1-yl)-6-(4-fluoro-phenyl) pteridine

25 The following illustrates the method step (d) shown in figure 4. The crude product obtained in example 54 was dissolved in a 1.25 M HCl solution in MeOH (20 ml) and 4-fluorophenylglyoxalmonoxime (504 mg, 3.0 mmole) was added portionwise. The mixture was refluxed for 3 hours. The reaction mixture was cooled down to room temperature and neutralized with NH₃ 25 % aqueous solution to pH 9. The yellow precipitate was filtered off and washed with water. The precipitate was adsorbed on silica gel and purified by flash chromatography, the mobile phase consisting of CH₃OH/CH₂Cl₂ mixtures (in a ratio gradually
30 ranging from 1:99 to 4:96), thus affording the title compound as a yellow powder (734 mg, 53 % yield), which was characterized by its mass spectrum MS as follows: MS *m/z* (%): 368 ([M+H]⁺, 100).

Example 58 - synthesis of 2-amino-4-(N-acetylpiperazin-1-yl)-6-(4-chloro-phenyl) pteridine

35 The method of example 57 was repeated, except for using 4-chlorophenylglyoxalmonoxime (554 mg, 3.0 mmole) instead of 4-fluorophenylglyoxalmonoxime. This afforded the title compound as a yellow powder (924 mg, 64 % yield), which was characterized by its mass spectrum MS as follows: MS *m/z* (%): 384 ([M+H]⁺, 100).

Example 59 - synthesis of 2-amino-4-[N-acetylpiperazin-1-yl]-6-(4-acetamido-phenyl) pteridine

The method of example 57 was repeated, except for using 4-acetyl-benzamidophenylglyoxalmonoxime (206 mg, 3.0 mmole) instead of 4-fluoro-phenylglyoxalmonoxime, and performing silica gel flash chromatography with a CH₃OH/CH₂Cl₂ gradient from 2:98 to 10:90. This afforded the title compound as a yellow powder (871 mg, 57 % yield), which was characterized by its mass spectrum MS as follows: MS *m/z* (%): 407 ([M+H]⁺, 100).

Examples 60 to 63 - synthesis of 2-amino-4-[N-(α -aminoacyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridines

The following illustrates the method step (a) shown in figure 5, wherein R₁₁ is a radical having an amino group in α position with respect to the carboxylic acid group. The compound of example 35 (0.367 g, 1 mmole) and a tert-butoxycarbonyl-protected amino-acid, such as L-phenylalanine (example 60), L-tyrosine (example 61), L-proline (example 62) or L-tryptophane (example 63), were suspended in dry dimethylformamide at room temperature under nitrogen and diisopropylethylamine (0.418 ml, 2.4 mmole), followed by *o*-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium tetrafluoroborate (0.482 g, 1.5 mmole) were added. The mixture was stirred for 2 hours and diluted with dichloromethane (50 ml). The organic layer was washed with a saturated solution of sodium hydrogen carbonate (50 ml), dried over anhydrous sodium sulfate and evaporated to dryness. The crude residue was purified by silica gel chromatography, the mobile phase consisting of CH₃OH/CH₂Cl₂ mixtures (in a ratio gradually ranging from 3:97 to 6:94), with 0.5% concentrated aqueous ammonia if needed. This procedure provided tert-butoxycarbonyl-protected 2-amino-4-[N-(α -aminoacyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine intermediates with yields ranging from 50 % to 80 % depending upon the tert-butoxycarbonyl-protected amino-acid used.

Then the said tert-butoxycarbonyl-protected intermediate (0.5 mmole) was deprotected either by being suspended in a mixture of dioxane (10 ml) and HCl 6M (20 ml) and stirred at room temperature until complete mixture or by using a solution of 20 % trifluoroacetic acid in dichloromethane (10 ml). The medium treated with HCl was then neutralized with NaOH 10M and volatiles were removed, whereas the mixture treated with trifluoroacetic acid was directly evaporated to dryness. The residue was adsorbed on silica and purified by silica gel column chromatography, the mobile phase consisting of CH₃OH/CH₂Cl₂ mixtures (in a ratio gradually ranging from 4:96 to 6:94) containing 0.5% of concentrated aqueous ammonia.

This procedure provided, with a yield ranging from 50 % to 70 % depending upon the tert-butoxycarbonyl-protected amino-acid used, the following pure final pteridine derivatives as yellow powders which were characterized by their mass spectrum MS as follows:

- 2-amino-4-[N-(2-(S)-amino-3-phenylpropionyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 60) : MS 515 ($[M+H]^+$;
- 2-amino-4-[N-(2-(S)-amino-3-(4-hydroxyphenyl)propionyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 61) : MS 531 ($[M+H]^+$;
- 5 - 2-amino-4-[N-(pyrrolidin-2-(S)-yl)carbonyl-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 62) : MS 465 ($[M+H]^+$; and
- 2-amino-4-[N-(2-(S)-amino-3-(indol-2-yl)propionyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 63) : MS 554 ($[M+H]^+$.

10 Example 64 - synthesis of 2-amino-4-(N-phenylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine

To a suspension of the compound of example 6 (196 mg, 0.5 mmole) in dioxane (10 ml) was added N-phenylpiperazine (0.23 ml, 1.5 mmole). The suspension was stirred at room temperature overnight. The precipitate was filtered off and washed with dioxane and
 15 diethylether, yielding the crude 2-acetylamino-4-(N-phenylpiperazine)-6-(3,4-dimethoxyphenyl)pteridine. Deprotection of the acetylamino group was achieved by dissolving this crude compound in methanol (5 ml) and a 20 % K_2CO_3 solution in water (5 ml). The solution was stirred overnight. Solvents were evaporated *in vacuo* and the residue was purified by preparative TLC (silica, using a CH_3OH/CH_2Cl_2 (5:95) mixture as an eluent),
 20 affording the title compound as a yellow powder (84 mg, yield 38 %) which was characterized by its mass spectrum MS as follows: MS *m/z* (%): 444 ($[M+H]^+$, 100).

Example 65 - synthesis of 2-amino-4-(N-benzylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine

25 Repeating the method of example 64, except for using N-benzyl-piperazine (0.26 ml, 1.5 mmole) instead of N-phenylpiperazine, afforded the title compound as a yellow powder (75 mg, yield 33%) which was characterized by its mass spectrum MS as follows: MS *m/z* (%): 458 ($[M+H]^+$, 100).

30 Example 66 - synthesis of 2-amino-4-(N-trans-cinnamylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine

Repeating the method of example 64, except for using N-cinnamyl-piperazine (0.306 ml, 1.5 mmole) instead of N-phenylpiperazine, afforded the title compound as a yellow powder (99 mg, yield 41%) which was characterized by its mass spectrum MS as follows: MS *m/z*
 35 (%): 484 ($[M+H]^+$, 100).

Examples 67 to 70 - synthesis of 2-substituted 4,6-diamino-5-nitroso-pyrimidines

To a suspension of 4,6-diamino-2-methylmercapto-5-nitroso-pyrimidine (1 g, 5.41 mmole), which may be prepared and characterised for instance as disclosed by Baddiley et al.
 40 in J. Chem. Soc. (1943) 383, in water (25 ml) was added a large excess (162 mmole) of an

appropriate amine. After heating the reaction mixture at 65 °C during 3 hours, a pink suspension was formed. The reaction mixture was then cooled down to + 4°C for 4 days. The pink precipitate was filtered off and washed with water, yielding the pure following compounds, each being characterised by its mass spectrum (MS), in yields ranging from 30 to

5 50 % :

- 2-phenylethylamino-4,6-diamino-5-nitroso-pyrimidine (example 67) was obtained from phenylethylamine ; MS: m/z (%): 259 ($[M+H]^+$, 100).
- 2-(2-thienylmethylamino)-4,6-diamino-5-nitroso-pyrimidine (example 68) was obtained from 2-thiophenemethylamine ; MS: m/z (%): 251 ($[M+H]^+$, 100).
- 10 - 2-pyrrolidino-4,6-diamino-5-nitroso-pyrimidine (example 69) was obtained from pyrrolidine ; MS: m/z (%): 209 ($[M+H]^+$, 100).
- 2-benzylamino-4,6-diamino-5-nitroso-pyrimidine (example 70) was obtained from benzylamine ; MS: m/z (%): 245 ($[M+H]^+$, 100).

15 Example 71 to 74 - synthesis of 2-substituted-4,5,6-triamino-pyrimidine sulfates

To a suspension of a 2-substituted-4,6-diamino-5-nitroso-pyrimidine obtained in one of examples 67 to 70 (1 mmole) in water (25 ml) was added portionwise sodium dithionite (3 mmole). The resulting suspension was refluxed until a yellow solution was formed. A sulfuric acid solution (2.5 ml of a 50 % solution in water) was then added. The reaction mixture was

20 cooled down to + 4°C for 5 hours. The white precipitate formed was filtered off, yielding the pure following compounds in yields ranging from 60% to 75%.

- 2-phenylethylamino- 4,5,6-triamino-pyrimidine sulfate (example 71),
- 2-(2-thienylmethylamino)- 4,5,6-triamino-pyrimidine sulfate (example 72),
- 2-pyrrolidino- 4,5,6-triamino-pyrimidine sulfate (example 73), and
- 25 - 2-benzylamino- 4,5,6-triamino-pyrimidine sulfate (example 74).

Examples 75 to 78 - synthesis of 2-substituted-4,5,6-triamino-pyrimidine dihydrochlorides

To a suspension of a 2-substituted-4,5,6-triamino-pyrimidine sulfate obtained in one of examples 71 to 74 (1 mmole) in water (6 ml) at 80 °C was added dropwise a solution of

30 barium chloride dihydrate (0.9 mmole) in water (2 ml). The resulting suspension was stirred for 30 minutes at 80 °C, then the reaction mixture was cooled down and barium sulfate was filtered off over Celite. The filtrate was evaporated in vacuo and co-evaporated with toluene yielding each of the following compounds as a yellow powder in yields ranging from 90 % to 98 %:

- 35 - 2-phenylethylamino- 4,5,6-triamino-pyrimidine dihydrochloride (example 75),
- 2-(2-thienylmethylamino)- 4,5,6-triamino-pyrimidine dihydrochloride (example 76),
- 2-pyrrolidino- 4,5,6-triamino-pyrimidine dihydrochloride (example 77), and
- 2-benzylamino- 4,5,6-triamino-pyrimidine dihydrochloride (example 78).

Examples 79 to 82 - synthesis of 2-substituted-4-amino-6-(3,4-dimethoxy-phenyl) pteridines

The following procedure is in accordance with step (e) of figure 6. To a solution of a 2-substituted-4,5,6-triamino-pyrimidine dihydrochloride obtained in one of examples 75 to 78 (1 mmole) in methanol (15 ml) was added the 3,4-dimethoxyphenylglyoxaloxime obtained according to example 3 (1 mmole). The resulting solution was refluxed for 2 hours, thus forming a yellow suspension. The reaction mixture was cooled down and neutralised by addition of a 33% aqueous ammonia solution until pH 9 was reached. The yellow precipitate was filtered off and further purified by silica gel flash chromatography (eluting with solvent mixture CH₃OH/CH₂Cl₂, gradient from 1:99 to 3:97), yielding as a yellow powder each of the pure following compounds, which was characterised by its mass spectrum (MS) and its ultraviolet spectrum (UV):

- 2-phenylethylamino-4-amino-6-(3,4-dimethoxyphenyl) pteridine (example 79) was obtained from the salt of example 75; MS: m/z (%): 403 ([M+H]⁺, 100), 827 ([2M+Na]⁺, 20); UV (MeOH, nm): 287, 315, 412.
- 2-(2-thienylmethylamino)- 4-amino-6-(3,4-dimethoxyphenyl) pteridine (example 80) was obtained from the salt of example 76; MS: m/z (%): 394 ([M+H]⁺, 100); UV (MeOH, nm): 287, 314, 410.
- 2-pyrrolidino-4-amino-6-(3,4-dimethoxyphenyl) pteridine (example 81) was obtained from the salt of example 77; MS: m/z (%): 353 ([M+H]⁺, 100), 727 ([2M+Na]⁺, 10); UV (MeOH, nm): 319, 423.
- 2-benzylamino-4-amino-6-(3,4-dimethoxyphenyl) pteridine (example 82) was obtained from the salt of example 78; MS: m/z (%): 389 ([M+H]⁺, 100); UV (MeOH, nm): 287, 315, 411.

Example 83 - synthesis of 4,5,6-triamino-2-methylmercaptopyrimidine dihydrochloride

To a suspension of a 2-methylmercapto-4,5,6-triamino-pyrimidine sulfate (44.3 mmole), which may be prepared and characterised for instance as disclosed by Taylor et al. in J. Am. Chem. Soc. (1952) 74:1644-1647, in water (135 ml) at 80 °C was added dropwise a solution of barium chloride dihydrate (39.8 mmole) in water (25 ml). The suspension was stirred for 30 minutes at 80 °C. The reaction mixture was cooled down and barium sulfate was filtered off over Celite. The filtrate was evaporated in vacuo and co-evaporated with toluene yielding the title compound as a yellow powder (10.2 g, 94 % yield).

Example 84 - synthesis of 4-amino-2-methylmercapto-6-(3,4-dimethoxy-phenyl) pteridine

To a suspension of 4,5,6-triamino-2-methylmercaptopyrimidine dihydrochloride (7.42 mmole, 1.81 g) in methanol (20 ml) was added a solution of 3,4-dimethoxyphenylglyoxaloxime (5.94 mmole, 1.24 g) in methanol. The resulting reaction mixture was refluxed for 3 hours. The reaction mixture was neutralised with concentrated aqueous ammonia until pH 9 was reached. The resulting precipitate was filtered off and further purified by flash chromatography

(silica, using an ethyl acetate / hexane mixture in a 4:6 ratio) yielding the pure title compound as a yellow powder which was characterised as follows: MS: m/z (%) 330 ($[M+H]^+$, 100), 681 ($[2M+Na]^+$, 30); UV (MeOH, nm): 292, 397.

5 Example 85 - synthesis of 4-amino-2-methylmercapto-6-phenyl-pteridine

A method similar to that of example 84 was used, starting from phenylglyoxal monoxime instead of 3,4-dimethoxyphenylglyoxalmonoxime. The title compound was characterised as follows: MS: m/z (%): 270 ($[M+H]^+$, 100); UV (MeOH, nm): 286, 379.

10 Example 86 - synthesis of 4-amino-2-morpholino-6-(3,4-dimethoxyphenyl) pteridine

A solution of the compound of example 84 (100 mg, 0.304 mmole) in morpholine (12 ml) was refluxed overnight. The solvents were removed in vacuo and the residue was purified first by flash chromatography (silica, gradient from 2:98 to 3:97 CH₃OH /CH₂Cl₂) and then by preparative TLC (silica, EtOAc/hexane 1 :1) yielding the title compound as a yellow powder
15 (70 mg, 63 % yield) characterised as follows: MS: m/z (%): 369 ($[M+H]^+$, 100), 759 ($[2M+Na]^+$, 20); UV (MeOH, nm): 297, 315, 418.

Example 87 - synthesis of 4-amino-2-piperidino-6-(3,4-dimethoxyphenyl) pteridine

A method similar to that of example 86 was used, starting from piperidine instead of
20 morpholine. The title compound obtained as a yellow powder (58 mg, 52 %) was characterised as follows: MS: m/z (%): 367 ($[M+H]^+$, 100), 755 ($[2M+Na]^+$, 10); UV (MeOH, nm): 319, 425.

Example 88 - synthesis of 2-amino-4-(homopiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine

25 Homopiperazine (1.39 g) was added to a stirred suspension of 2-amino-6-(3,4-dimethoxyphenyl)pteridine (520 mg) in pyridine (9 ml) and 1,1,1,3,3,3-hexamethyldisilazane (9.2 ml) in the presence of a catalytic amount of ammonium sulfate (54 mg) and *p*-toluenesulfonic acid (52 mg). The mixture was heated under reflux for 72 hours until a clear solution was obtained. The mixture was cooled down and the solvents were evaporated *in vacuo*. The residue was adsorbed on silica and purified by silica gel column chromatography,
30 using a 9:1 CH₂Cl₂/CH₃OH mixture containing 1 % concentrated aqueous ammonia as eluent, affording the desired compound (305 mg, yield 46 %) which was characterized by its mass spectrum as follows : m/z (%) 785 ($[2M+H]^+$, 15), 382 ($[M+H]^+$, 100).

35 Example 89 - synthesis of 2-amino-4-(*N*-phenoxyacetyl)-homopiperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine

To a solution of 2-amino-4-(homopiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (160 mg) in DMF (20 ml) was added triethylamine (0.55 mmole) and phenoxyacetyl chloride (0.5 mmole). The solution was stirred at room temperature for 24 hours. The solution was

diluted with CH_2Cl_2 and extracted 3 times with water. The organic solvents were evaporated *in vacuo*. The residue was adsorbed on silica, and purified by flash chromatography (silica, the mobile phase being $\text{CH}_3\text{OH}/\text{CH}_2\text{Cl}_2$ mixtures (in a ratio gradually ranging from 2:98 to 5:95). This procedure provided with a yield of 67 % the title compound as a yellow powder (145 mg)

5 which was characterized as follows:

- mass spectrum : m/z (%) 1053 ($[\text{2M}+\text{H}]^+$, 5), 516 ($[\text{M}+\text{H}]^+$, 100), and
- UV spectrum (nm) : 213, 296, 408.

10 Examples 90 to 98 - synthesis of 2-amino-4-[N-(thio)carboxy]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl)pteridines

To a solution of 2-amino-4-(piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (200 mg, 0.55 mmole) in DMF (20 ml) was added triethylamine (0.65 mmole, 92 μl) and a suitable chloroformate (0.71 mmole). The solution was stirred at room temperature for 2 to 24 hours, depending upon the chloroformate used, while monitoring the reaction by TLC. The solution
15 was diluted with CH_2Cl_2 and extracted with water (3 times). The organic solvents were evaporated *in vacuo*. The residue was adsorbed on silica, and purified by silica gel column chromatography, the mobile phase being $\text{CH}_3\text{OH}/\text{CH}_2\text{Cl}_2$ mixtures (in a ratio gradually ranging from 2:98 to 5:95). This procedure provided with a yield ranging from 60 % to 80 %, depending on the chloroformate used, the following pure pteridine derivatives, which were
20 characterized by their mass spectrum (MS) and their ultraviolet spectrum (UV).

- 2-amino-4-[(N-4-methyl-phenoxy-carbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine, obtained from *p*-tolylchloroformate (example 90) : MS : m/z (%) 502 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 215, 296, 412 ;
- 2-amino-4-[(N-4-methoxy-phenoxy-carbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-
25 phenyl)pteridine, obtained from *p*-methoxy-phenyl chloroformate (example 91) : MS : m/z (%) 518 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 217, 296, 412 ;
- 2-amino-4-[(N-4-fluoro-phenoxy-carbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 92), obtained from *p*-fluoro-phenyl chloroformate : MS : m/z (%) 506 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 213, 296, 412 ;
- 2-amino-4-[N-(2-methoxy)-phenoxy-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl)
30 pteridine (example 93), obtained from 2-methoxy-phenylchloroformate : MS : m/z (%) 518 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 215, 296, 410 ;
- 2-amino-4-[N-(4-chloro)-phenoxy-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 94), obtained from *p*-chloro-phenyl chloroformate : MS : m/z (%) 523
35 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 217, 296, 412 ;
- 2-amino-4-[N-isobutoxy-carbonyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl)pteridine (example 95), obtained from isobutyl chloroformate : MS : m/z (%) 468 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 215, 297, 413 ;

- 2-amino-4-[N-(2-chloro)-phenoxy-carbonyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 96), obtained from 2-chloro-phenyl chloroformate : MS : m/z (%) 523 ([M+H]⁺, 100) ; UV (nm) : 213, 296, 412 ;
- 5 - 2-amino-4-[N-(2-methoxy)-ethoxy-carbonyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 97), obtained from 2-methoxy-ethyl chloroformate : MS : m/z (%) 470 ([M+H]⁺, 100) ; UV (nm) : 212, 256, 296, 412 ; and
- 2-amino-4-[N-(2-naphthoxy)-carbonyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 98), obtained from 2-naphthyl chloroformate : MS : m/z (%) 538 ([M+H]⁺, 100) ; UV (nm) : 222, 296, 412.

10

Examples 99 to 109 - synthesis of 2-amino-4-(N-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridines

To a solution of 2-amino-4-(piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (0.61 mmol, 225 mg) in DMF (30 ml) was added a suitable isocyanate (0.92 mmole). The solution
 15 was stirred at room temperature for 2 to 24 hours, depending upon the isocyanate used, the reaction being monitored by TLC. The solution was diluted with CH₂Cl₂ and extracted 3 times with water. The organic solvents were evaporated *in vacuo*. The residue was adsorbed on silica, and purified by silica gel column chromatography, the mobile phase being CH₃OH/CH₂Cl₂ mixtures (in a ratio gradually ranging from 2:98 to 5:95). This procedure
 20 provided with a yield ranging from 60% to 80%, depending on the isocyanate used, the following pure pteridine derivatives, each as a yellow powder, which were characterized by their mass spectrum (MS) and their ultraviolet spectrum (UV):

- 2-amino-4-(N-phenyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (example 99), obtained from phenyl isocyanate : MS : m/z (%) 487 ([M+H]⁺, 100) ; UV
 25 (nm) : 239, 297, 412 ;
- 2-amino-4-[N-4-fluorophenyl-carbamoyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl)pteridine (example 100), obtained from 4-fluoro-phenyl isocyanate : MS : m/z (%) 1031 ([2M+Na]⁺, 15), 523 ([M+H]⁺, 100) ; UV (nm) : 297, 413 ;
- 2-amino-4-(N-4-methylphenyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 101), obtained from 4-methyl-phenyl isocyanate : MS : m/z (%) 1023
 30 ([2M+Na]⁺, 15), 501 ([M+H]⁺, 100) ; UV (nm) : 241, 270, 413 ;
- 2-amino-4-(N-4-cyanophenyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (example 102), obtained from 4-cyano-phenyl isocyanate : MS : m/z (%) 512 ([M+H]⁺, 100) ;
- 35 - 2-amino-4-(N-3-methylphenyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (example 103), obtained from 3-methyl-phenyl isocyanate : MS : m/z (%) 501 ([M+H]⁺, 100) ; UV (nm) : 241, 297, 412 ;

- 2-amino-4-(*N*-benzylcarbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (example 104), obtained from benzyl isocyanate : MS : m/z (%) 501 ($[M+H]^+$, 100) ; UV (nm) : 242, 297, 413 ;
- 2-amino-4-(*N*-4-fluorobenzylcarbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (example 105), obtained from 4-fluorobenzyl isocyanate : MS : m/z (%) 1059 ($[2M+Na]^+$, 10), 519 ($[M+H]^+$, 100) ; UV (nm) : 212, 297, 412 ;
- 2-amino-4-(*N*-3-chloro-4-fluorophenylcarbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridine (example 106), obtained from 3-chloro-4-fluoro-phenyl isocyanate : MS : m/z (%) 540 ($[M+H]^+$, 100) ; UV (nm) : 212, 240, 296, 412 ;
- 2-amino-4-(*N*-3-thienylcarbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl)pteridines (example 107), obtained from 3-thienyl isocyanate : MS : m/z (%) 493 ($[M+H]^+$, 100) ; UV (nm) : 216, 297, 413 ;
- 2-amino-4-[*N*-2-(2-thienyl)ethylcarbamoyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl)pteridine (example 108), obtained from 2-(2-thienyl)ethyl isocyanate ; and
- 2-amino-4-[(*N*-butyl-carbamoyl-piperazin-1-yl)]-6-(3,4-dimethoxyphenyl)pteridine (example 109), obtained from *n*-butyl isocyanate : MS : m/z (%) 467 ($[M+H]^+$, 100) ; UV (nm) : 214, 298, 413.

20 Examples 110 and 111 - synthesis of 2-amino-4-[*N*-(α -aminoacyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridines

The procedure of examples 60 to 63 was repeated while starting from different *tert*-butoxycarbonyl-protected amino-acids, i.e. glycine (example 110) and L-asparagine (example 111). The procedure provided the two following pure pteridine derivatives as yellow powders which were characterized by their mass spectrum MS as follows :

- 2-amino-4-[*N*-aminoacetyl]-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 110): MS m/z (%) 425 ($[M+H]^+$); and
- 2-amino-4-[*N*-[2-(*S*),4-diamino-4-oxobutanoyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 111): MS m/z (%) 482 ($[M+H]^+$, 100).

30 Examples 112 to 115 - synthesis of 2-amino-4-(*N*-acyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridines

- 35 The compound of example 35 (0.367 g, 1 mmole) and a carboxylic acid or anhydride such as mono-methyl terephthalate (example 112), dimethylglycine (example 113), succinamic acid (example 114) or succinic anhydride (example 115) were suspended in dry DMF at room temperature under a nitrogen atmosphere and then diisopropylamine (0.418 ml, 2.4 mmole), followed by *o*-benzotriazol-1-yl-*N,N,N',N'*-tetramethyluronium tetrafluoroborate (0.482 mg, 1.5 mmole) were added. The mixture was stirred until completion of the reaction and then diluted with dichloromethane (50 ml). The organic layer was washed with a saturated solution of sodium hydrogenocarbonate (50 ml), dried over anhydrous sodium sulfate and

evaporated to dryness. The crude residue was purified by silica gel column chromatography, the mobile phase consisting of CH₃OH/CH₂Cl₂ mixtures (in a ratio gradually ranging from 2:98 to 10:90), with 0.5 % concentrated ammonia or acetic acid if needed. This procedure provided, with yields ranging from 56 % to 72 % depending upon the starting carboxylic acid or anhydride, the desired compounds which were characterized by their mass spectrum MS as follows :

- 2-amino-4-[N-[4-(methoxycarbonyl)benzoyl]-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 112): MS m/z (%) 530 ([M+H]⁺, 100);
- 2-amino-4-[N-[4-(dimethylamino)acetyl]-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 113): MS m/z (%) 453 ([M+H]⁺, 100);
- 2-amino-4-[N-(4-amino-4-oxo-butanoil)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 114): MS m/z (%) 467 ([M+H]⁺, 100) ; and
- 2-amino-4-[N-(3-carboxypropanoal)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine (example 115): MS m/z (%) 468 ([M+H]⁺, 100).

Example 116 - synthesis of 2-amino-4-[N-[4-(carboxy)benzoyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine

2-amino-4-[N-[4-(methoxycarbonyl)benzoyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (0.212 g, 1.4 mmole) was dissolved in THF (8 ml) and aqueous LiOH 0.1 N was added (8 ml). The mixture was stirred 24 hours at room temperature and the pH was adjusted at 3 with HCl 1 N. The precipitate was filtered, washed with H₂O, EtOAc, Et₂O and dried in a vacuum dessicator over P₂O₅, yielding the title compound as a yellow powder which was characterized by its mass spectrum: MS m/z (%) 516 ([M+H]⁺, 100).

Examples 117 to 119 - synthesis of 2-amino-4-[(N-alkyl-N-aryl)-carbamoyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl)pteridines

The synthesis of such compounds is achieved by the three-step procedure shown in figure 9, which is derived from the teaching of Batey et al. in *Tetrahedron Lett.* (1998) 39 : 6267-6270. The detailed procedure is as follows :

- (a) to a suspension of carbonyl diimidazole (30.4 mmole, 4.93 g) in THF (50 ml) was added a suitable *N*-alkyl-aniline derivative (28 mmole), such as for example *N*-methylaniline (example 117), *N*-ethylaniline (example 118) or *N*-methyl-*p*-toluidine (example 119). The mixture was refluxed for 24 hours, after which an additional amount of carbonyl diimidazole (2.24 g) was added. The reaction mixture was refluxed for another 6 hours, until the reaction reaches completion (TLC monitoring). After cooling down the reaction mixture, the solvents were evaporated *in vacuo* yielding the crude *N*-alkyl-aniline carbamoyl imidazoles which were used in the next step without any further purification.
- (b) to a solution of the crude *N*-alkyl-aniline carbamoyl imidazole (32 mmole) in acetonitrile (50 ml) was added methyl iodide (128 mmole). The mixture was stirred at room

temperature for 24 hours. The solvent was evaporated *in vacuo* yielding N-alkyl-aniline carbamoyl N-methyl-imidazolium iodide.

- (c) to a solution of 2-amino-4-(piperazin-1-yl)-6-(3,4-dimethoxy-phenyl)pteridine (224 mg, 0.61 mmole) in DMF (30 ml) was added triethylamine (111 μ l, 0.80 mmole) and a suitable N-alkyl-aniline carbamoyl N-methyl-imidazolium iodide (0.92 mmole). The reaction mixture was stirred at room temperature for 24 hours. The reaction was diluted with CH_2Cl_2 and extracted 3 times with H_2O . Evaporation of the solvents *in vacuo*, followed by purification of the residue by silica gel column chromatography, the mobile phase being $\text{CH}_3\text{OH}/\text{CH}_2\text{Cl}_2$ mixtures (in a ratio gradually ranging from 2:98 to 3:97) provided each of the title compounds as a yellow powder, with a yield ranging from 65 to 80 %, depending on the N-alkyl-aniline derivative used.

The following compounds were synthesized following this procedure and characterised by their mass spectrum (MS) and ultraviolet spectrum (UV) as follows:

- 2-amino-4-(N-methyl-N-phenyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 117) : MS : m/z (%) 501 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 213, 298, 412 ;
- 2-amino-4-(N-ethyl-N-phenyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 118) : MS : m/z (%) 515 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 213, 298, 413 ; and
- 2-amino-4-(N-methyl-N-tolyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 119) : MS : m/z (%) 515 ($[\text{M}+\text{H}]^+$, 100) ; UV (nm) : 213, 298, 412.

Example 120 – model of rheumatoid arthritis

Collagen type II (hereinafter referred as CII) induced experimental model of rheumatoid arthritis (hereinafter referred as RA) in DBA mice is widely accepted as the most relevant and predictive preclinical model for RA. In this model, DBA mice are immunized with CII, the collagen type mainly present in the joint structures, together with complete Freund Adjuvant in their tail. 2 to 3 weeks later, several of the immunized mice start to develop arthritis in the four footpaths. In order to further worsen the disease, mice are given a second CII boost at three weeks after the first immunisation, this time however in a footpath. Because the immune system is already immunised in these mice, this rapidly provokes a severe swelling of the injected footpath (named Delayed Type Hypersensitivity or DTH) which can be used as a measurement for T-cell activation. Within a few days after the booster, almost all untreated animals start developing symptoms of arthritis. RA development is scored from 0 to 16 (16 being severe clinical arthritis in all four footpaths). At the end of the study (3 weeks after the CII boost) antibody formation was determined against CII and histology performed on the footpaths.

The efficiency of the pteridine derivative of example 17 (administered in an amount of 20 mg/kg/day, started one day before the CII boost) was explored in this CII-model. All such treated animals developed significantly less severe rheumatoid arthritis (clinical scores ranging from 2 to 4), as compared to untreated control mice (clinical scores ranging from 6 to

12) and also compared to mice treated with methotrexate (clinical scores ranging from 2 to 7), the most effective compound for the treatment of RA to date.

Additionally, increasing the dose of the pteridine derivative of example 17 up to 40 mg/kg/day has no mortality or cytotoxic effect on mice *in vivo*, whereas increasing the dose of methotrexate (10 mg/kg/day, 3 times a week) leads to death of all animals.

Example 121 – model of protection against septic shock

As a control group, 4 sham treated (saline injection) C3H mice being injected intraperitoneously with 100 µg lipopolysaccharide (hereinafter LPS) per mouse, all died within 1 to 3 days after injection. However, when four C3H mice being injected intraperitoneously with 100 µg lipopolysaccharide (hereinafter LPS) per mouse were treated during 2 days with the pteridine derivative of example 17 (one first intraperitoneous injection of 20 mg/kg/day at the time of injection, and a second injection 24 hours later), all mice were protected from acute shock related mortality.

Examples 122 to 162 - synthesis of 2-amino-4-(N-substituted-piperazino)-6-(3,4-dimethoxyphenyl) pteridines

The following procedure is similar to that of examples 64 to 66. To a suspension of the compound of example 6 (1 mmole) in dioxane (20 ml) was added a suitable N-substituted piperazine (1.5 mmole). The suspension was stirred at room temperature for 16 hours. The solvents were evaporated in vacuo yielding crude 2-acetylamino-4-(N-substituted piperazino)-6-(3,4-dimethoxyphenyl) pteridine. Deprotection of the 2-acetylamino group was achieved by dissolving this crude compound in 20 ml of a 1:1 mixture of methanol and 20% K₂CO₃ in water. The solution was stirred for 16 hours at room temperature. Solvents were evaporated in vacuo and the residue was purified by preparative TLC (silica, using a CH₃OH/CH₂Cl₂ (5:95) mixture as an eluent), affording the following compounds as yellow powders in yields ranging from 20 to 70%:

- 2-amino-4-(1-(2-methoxyethyl)piperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 122) was obtained from 1-(2-methoxyethyl) piperazine and characterised as follows: MS: m/z (%): 873 ([2M+Na]⁺, 15), 426 ([M+H]⁺, 100);
- 2-amino-4-(1-cyclohexylmethyl)piperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 123) was obtained from 1-(cyclohexylmethyl)piperazine and characterised as follows: MS: m/z (%): 949 ([2M+Na]⁺, 5), 464 ([M+H]⁺, 100);
- 2-amino-4-(1-cyclopentylpiperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 124) was obtained from 1-cyclopentylpiperazine and characterised as follows: MS: m/z (%): 893 ([2M+Na]⁺, 25), 436 ([M+H]⁺, 100]; UV (MeOH, nm): 213, 296, 413;
- 2-amino-4-(1-butylpiperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 125) was obtained from 1-(butyl)piperazine and characterised as follows: MS: m/z (%): 893 ([2M+Na]⁺, 25), 436 ([M+H]⁺, 100]; UV (MeOH, nm): 216, 295, 413;

- 2-amino-4-(1-isopropylpiperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 126) was obtained from 1-(isopropyl) piperazine and characterised as follows: MS: m/z (%): 841 ([2M+Na]⁺, 20), 410 ([M+H]⁺, 100); UV (MeOH, nm): 215, 295, 412;
- 5 - 2-amino-4-(1-(2-diethylaminoethyl)-piperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 127) was obtained from 1-(2-diethylaminoethyl)-piperazine and characterised as follows: MS: m/z (%): 955 ([2M+Na]⁺, 20), 437 ([M+H]⁺, 100); UV (MeOH, nm): 216, 297, 413;
- 10 - 2-amino-4-(1-(2-diisopropylaminoethyl)-piperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 128) was obtained from 1-(2-diisopropylaminoethyl)-piperazine and characterised as follows: MS: m/z (%): 495 ([M+H]⁺, 100); UV (MeOH, nm): 215, 297, 413;
- 2-amino-4-(1-(2-morpholino-4-yl-ethyl)-piperazino)-6-(3,4-dimethoxyphenyl) pteridine (example 129) was obtained from 1-(2-morpholino-4-yl-ethyl)-piperazine and characterised as follows: MS m/z (%): 481 ([M+H]⁺, 100); UV (MeOH, nm): 217, 297, 15 414;
- 2-amino-4-(4-[2-(piperazin-1-yl)-acetyl]-morpholino)-6-(3,4-dimethoxy-phenyl) pteridine (example 130) was obtained from 4-[2-(piperazin-1-yl)-acetyl] morpholine and characterised as follows: MS: m/z (%): 495 ([M+H]⁺, 100); UV (MeOH, nm): 219, 297, 414;
- 20 - 2-amino-4-(4-[2-(piperazin-1-yl)-acetyl]-pyrrolidino)-6-(3,4-dimethoxy-phenyl) pteridine (example 131) was obtained from 4-[2-(piperazin-1-yl)-acetyl] pyrrolidine and characterised as follows: MS: m/z (%): 979 ([2M+Na]⁺, 20), 479 ([M+H]⁺, 100); UV (MeOH, nm): 219, 307, 411;
- 2-amino-4-(2-[piperazin-1-yl]-acetic acid N-methyl N-phenyl amide)-6-(3,4-dimethoxyphenyl) pteridine (example 132) was obtained from 2-[piperazin-1-yl]-acetic acid N-methyl-N-phenylamide and characterised as follows: MS: m/z (%): 515 ([M+H]⁺, 100); UV (MeOH, nm): 219, 307, 411;
- 25 - 2-amino-4-(2-(piperazin-1-yl)-propionic acid ethyl ester)-6-(3,4-dimethoxyphenyl) pteridine (example 133) was obtained from 2-(piperazin-1-yl)-propionic acid ethyl ester (in order to avoid transesterification, a mixture of ethanol and sodium was used for the deprotection of the acetyl group) and characterised as follows: MS: m/z (%): 468 ([M+H]⁺, 100); UV (MeOH, nm): 216, 297, 413;
- 30 - 2-amino-4-(3-(piperazin-1-yl)-propionic acid ethyl ester)-6-(3,4-dimethoxyphenyl) pteridine (example 134) was obtained from 3-(piperazin-1-yl)-propionic acid ethyl ester (in order to avoid transesterification, a mixture of ethanol and sodium (15 equivalents) was used for the deprotection of the acetyl group) and characterised as follows: MS: m/z (%): 957 ([2M+Na]⁺, 10), 468 ([M+H]⁺, 100); UV (MeOH, nm): 216, 296, 412;
- 35 - 2-amino-4-(2-(piperazin-1-yl)-acetic acid ethyl ester)-6-(3,4-dimethoxyphenyl) pteridine (example 135) was obtained from 3-(piperazin-1-yl)-propionic acid ethyl ester (in order to

- avoid transesterification, a mixture of ethanol and sodium (15 equivalents) was used for the deprotection of the acetyl group) and characterised as follows: MS: m/z (%): 454 ($[M+H]^+$, 100); UV (MeOH, nm): 216, 297, 413;
- 2-amino-4-(1-(3-methyl-benzyl)piperazinyl)-6-(3,4-dimethoxyphenyl) pteridine (example 136) was obtained from 1-(3-methylbenzyl)-piperazine and characterised as follows: MS: m/z (%): 965 ($[2M+Na]^+$, 10), 472 ($[M+H]^+$, 100);
 - 2-amino-4-[(2,6-dichlorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 137) was obtained from 1-(2,6-dichloro-benzyl)-piperazine and characterised as follows: MS: m/z (%): 526 ($[M+H]^+$, 100);
 - 2-amino-4-[(4-fluorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 138) was obtained from 1-(4-fluorobenzyl)-piperazine and characterised as follows: MS: m/z (%): 476 ($[M+H]^+$, 100);
 - 2-amino-4-[(4-chlorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 139) was obtained from 1-(4-chloro-benzyl)-piperazine and characterised as follows: MS: m/z (%): 492 ($[M+H]^+$, 100);
 - 2-amino-4-[(4-methylbenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 140) was obtained from 1-(4-methyl-benzyl)-piperazine and characterised as follows: MS: m/z (%): 472 ($[M+H]^+$, 100);
 - 2-amino-4-[(2-fluorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 141) was obtained from 1-(2-fluorobenzyl)-piperazine and characterised as follows: MS: m/z (%): 476 ($[M+H]^+$, 100);
 - 2-amino-4-[(3,4-dichlorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 142) was obtained from 1-(3,4-dichlorobenzyl)-piperazine and characterised as follows: MS: m/z (%): 526 ($[M+H]^+$, 100);
 - 2-amino-4-(piperonyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 143) was obtained from 1-piperonyl-piperazine and characterised as follows: MS: m/z (%): 502 ($[M+H]^+$, 100);
 - 2-amino-4-[(4-*tert*-butylbenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 144) was obtained from 1-(4-*tert*-butyl-benzyl) piperazine) and characterised as follows: MS: m/z (%): 514 ($[M+H]^+$, 100);
 - 2-amino-4-[(4-pyridyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 145) was obtained from 1-(4-pyridyl)-piperazine and characterised as follows: MS: m/z (%): 445 ($[M+H]^+$, 100); UV (MeOH, nm): 213, 289, 411;
 - 2-amino-4-[(2-pyridyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 146) was obtained from 1-(2-pyridyl)-piperazine and characterised as follows: MS: m/z (%): 911 ($[2M+Na]^+$, 60), 889 ($[2M+H]^+$, 60), 445 ($[M+H]^+$, 100); UV (MeOH, nm): 215, 247, 297, 413;

- 2-amino-4-((2-pyrimidinyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 147) was obtained from 1-(2-pyrimidinyl)-piperazine and characterised as follows: MS: m/z (%): 446 ($[M+H]^+$, 100); UV (MeOH, nm): 216, 244, 297, 413;
- 5 - 2-amino-4-((3-methoxyphenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 148) was obtained from 1-(3-methoxyphenyl)-piperazine and characterised as follows: MS: m/z (%): 969 ($[2M+Na]^+$, 15), 446 ($[M+H]^+$, 100); UV (MeOH, nm): 215, 295, 413;
- 2-amino-4-(1-(3-phenylpropyl)-piperazine)-6-(3,4-dimethoxyphenyl) pteridine (example 149) was obtained from 1-(3-phenylpropyl)-piperazine and characterised as follows: MS: m/z (%): 486 ($[M+H]^+$, 100); UV (MeOH, nm): 217, 296, 413;
- 10 - 2-amino-4-((3,4-dichlorophenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 150) was obtained from 1-(3,4-dichlorophenyl)-piperazine and characterised as follows: MS: m/z (%): 512 ($[M+H]^+$, 100); UV (MeOH, nm): 213, 263, 295, 412;
- 2-amino-4-((3-dichlorophenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 151) was obtained from 1-(3-dichlorophenyl)-piperazine and characterised as follows: MS: m/z (%): 478 ($[M+H]^+$, 100); UV (MeOH, nm): 213, 257, 296, 413;
- 15 - 2-amino-4-((1-phenylethyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 152) was obtained from 1-(1-phenylethyl)-piperazine and characterised as follows: MS: m/z (%): 965 ($[2M+Na]^+$, 10), 472 ($[M+H]^+$, 100); UV (MeOH, nm): 216, 298, 413;
- 20 - 2-amino-4-((2-(1-pyrrolyl)-ethyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 153) was obtained from 1-[2-(1-(pyrrolyl)-ethyl)]-piperazine and characterised as follows: MS: m/z (%): 461 ($[M+H]^+$, 100); UV (MeOH, nm): 216, 297, 413;
- 2-amino-4-((2-phenoxyethyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 154) was obtained from 1-(2-phenoxyethyl)-piperazine and characterised as follows: MS: m/z (%): 488 ($[M+H]^+$, 100); UV (MeOH, nm): 216, 297, 413;
- 25 - 2-amino-4-(1-(2-imidazol-1-yl-ethyl)-piperazine)-6-(3,4-dimethoxyphenyl) pteridine (example 155) was obtained from 1-(2-imidazol-1-yl-ethyl)-piperazine and characterised as follows: MS: m/z (%): 945 ($[2M+Na]^+$, 10), 462 ($[M+H]^+$, 100); UV (MeOH, nm): 216, 297, 413;
- 30 - 2-amino-4-((3-pyridyl)-methyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 156) was obtained from 1-(3-pyridyl)-methyl-piperazine and characterised as follows: MS: m/z (%): 939 ($[2M+Na]^+$, 15), 459 ($[M+H]^+$, 100); UV (MeOH, nm): 215, 297, 413;
- 2-amino-4-((4-pyridyl)-methyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 157) was obtained from 1-(4-pyridyl)-methyl-piperazine and characterised as follows: MS: m/z (%): 939 ($[2M+Na]^+$, 15), 459 ($[M+H]^+$, 100); UV (MeOH, nm): 218, 297, 414;
- 35

- 2-amino-4-((1-naphthylmethyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 158) was obtained from 1-(1-naphthylmethyl-piperazine and characterised as follows: MS: m/z (%): 508 ($[M+H]^+$, 100); UV (MeOH, nm): 223, 297, 413;
- 2-amino-4-(*N*-phenethylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 159) was obtained from *N*-phenethylpiperazine and characterised as follows: MS: m/z (%): 965 ($[2M+Na]^+$, 10), 472 ($[M+H]^+$, 100); UV (MeOH, nm): 215, 297, 413;
- 2-amino-4-((2-methoxyphenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 160) was obtained from 1-(2-methoxyphenyl)-piperazine and characterised as follows: MS: m/z (%): 474 ($[M+H]^+$, 100); UV (MeOH, nm): 213, 295, 413;
- 2-amino-4-((4-methoxyphenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 161) was obtained from 1-(4-methoxyphenyl)-piperazine and characterised as follows: MS: m/z (%): 474 ($[M+H]^+$, 100); UV (MeOH, nm): 212, 296, 413; and
- 2-amino-4-((4-chlorophenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 162) was obtained from 1-(4-chlorophenyl)piperazine and characterised as follows: MS: m/z (%): 478 ($[M+H]^+$, 100); UV (MeOH, nm): 258, 296, 413.

Examples 163 to 180 - synthesis of 2-amino-6-(3,4-dimethoxyphenyl)-4-(substituted piperazin-1-yl) pteridines

The following procedure is in accordance with the scheme shown in figure 3. A mixture of the compound of example 4 (299 mg, 1.0 mmole), 1,1,1,3,3,3-hexamethyldisilazane (1 ml, 4.7 mmole), a *N*-substituted piperazine (4.0 mmole), *p*-toluenesulfonic acid (20 mg, 0.1 mmole) and ammonium sulfate (20 mg, 0.15 mmole) in toluene (4 ml) was refluxed for 48 hours (the reaction mixture became clear when the reaction was finished). After removing the solvents under reduced pressure, the residue was purified by flash chromatography over silica (CH₃OH/CH₂Cl₂ 1:20 to 1:30) reaching the desired following compounds as yellow solids in yields indicated below:

- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(3-propionitril)-piperazin-1-yl) pteridine (example 163) was obtained from 3-(1-piperazinyl)-propionitrile in 43 % yield and characterized as follows: R_f = 0.50 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 215, 297, 412; MS (m/z): 421 ($[M+H]^+$, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-(1,3)-dioxolan-2-yl-ethyl)-piperazin-1-yl) pteridine (example 164) was obtained from 2-[2-(piperazin-1-yl)-ethyl]-1,3-dioxolane in 55 % yield and characterized as follows: R_f = 0.49 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 215, 295, 412; MS (m/z): 468 ($[M+H]^+$, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-ethoxyethyl)-piperazin-1-yl) pteridine (example 165) was obtained from 1-(2-ethoxyethyl)-piperazine in 35 % yield and characterized as follows: R_f = 0.33 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 213, 295, 412; MS (m/z): 440 ($[M+H]^+$, 100);

- 2-amino-6-(3,4-dimethoxyphenyl)-4-(pent-3-yl-piperazin-1-yl) pteridine (example 166) was obtained from 1-(3-pentyl)-piperazine in 22 % yield and characterized as follows: Rf = 0.43 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 212, 293, 412; MS (m/z): 438.2 ([M+H]⁺, 100);
- 5 - 2-amino-6-(3,4-dimethoxyphenyl)-4-(1-pentyl-piperazin-1-yl) pteridine (example 167) was obtained from 1-(1-pentyl)-piperazine in 22 % yield and characterized as follows: Rf = 0.54 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 212, 294, 412; MS (m/z): 438 ([M+H]⁺, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(1-isobutyl-piperazin-1-yl) pteridine (example 168) was obtained from 1-isobutylpiperazine in 26 % yield and characterized as follows: Rf = 0.42 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 213, 294, 412; MS (m/z): 424 ([M+H]⁺, 100);
- 10 - 2-amino-6-(3,4-dimethoxyphenyl)-4-((tetrahydrofurfuryl)-piperazin-1-yl) pteridine (example 169) was obtained from 1-tetrahydrofurfuryl-piperazine in 31 % yield and characterized as follows: Rf = 0.37 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 215, 295, 412; MS (m/z): 453 ([M+H]⁺, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(1,3-dioxolan-2-yl-methylpiperazin-1-yl) pteridine (example 170) was obtained from 2-(piperazin-1-yl-methyl)-1,3-dioxolane in 65 % yield and characterized as follows: Rf = 0.46 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 217, 297, 413; MS (m/z): 454 ([M+H]⁺, 100);
- 15 - 2-amino-4-((3,5-dichlorophenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine (example 171) was obtained from 1-(3,5-dichlorophenyl)-piperazine in 83 % yield and characterised as follows: Rf = 0.70 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 216, 263, 296, 413; MS (m/z): 512.2 ([M+H]⁺, 100);
- 25 - 2-amino-6-(3,4-dimethoxyphenyl)-4-((4-fluorophenyl)-piperazin-1-yl) pteridine (example 172) was obtained from 1-(4-fluorophenyl)-piperazine in 20 % yield and characterised as follows: Rf = 0.53 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 212, 296, 413; MS (m/z): 462.2 ([M+H]⁺, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((3-trifluoromethylphenyl)-piperazin-1-yl) pteridine (example 173) was obtained from 1-(3-trifluoromethylphenyl)-piperazine in 71 % yield and characterised as follows: Rf = 0.58 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 212, 257, 296, 413; MS (m/z): 512.2 ([M+H]⁺, 100);
- 30 - 2-amino-6-(3,4-dimethoxyphenyl)-4-((3,4-dimethylphenyl)-piperazin-1-yl) pteridine (example 174) was obtained from 1-(3,4-dimethylphenyl)-piperazine in 48 % yield and characterised as follows: Rf = 0.58 (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 243, 296, 413; MS (m/z): 472.3 ([M+H]⁺, 100);
- 35 - 2-amino-6-(3,4-dimethoxyphenyl)-4-((3-methylphenyl)-piperazin-1-yl) pteridine (example 175) was obtained from 1-(3-methylphenyl)-piperazine in 59 % yield and characterised as follows:

follows: $R_f = 0.47$ (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 218, 244, 297, 413; MS (m/z): 458.2 ([M+H]⁺, 100);

- 2-amino-6-(3,4-dimethoxyphenyl)-4-((4-methylphenyl)-piperazin-1-yl) pteridine (example 176) was obtained from 1-(4-methylphenyl)-piperazine in 59 % yield and characterised as follows: $R_f = 0.50$ (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 213, 242, 296, 413; MS (m/z): 458.2 ([M+H]⁺, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((2-pyridyl)methyl-piperazin-1-yl) pteridine (example 177) was obtained from 1-((2-pyridyl)-methyl)-piperazine in 27 % yield and characterised as follows: $R_f = 0.38$ (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 216, 297, 413; MS (m/z): 459.2 ([M+H]⁺, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(thiazol-2-yl)-piperazin-1-yl) pteridine (example 178) was obtained from 1-thiazol-2-yl-piperazine in 11 % yield and characterised as follows: $R_f = 0.55$ (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 213, 294, 413; MS (m/z): 451.2 ([M+H]⁺, 100);
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(1-(1-methyl-piperidin-3-yl-methyl)-piperazin-1-yl) pteridine (example 179) was obtained from 1-(1-methyl-piperidin-3-yl-methyl)-piperazine in 48 % yield and characterised as follows: MS (m/z): 479 ([M+H]⁺, 100); UV (MeOH/H₂O, nm): 217, 266, 297, 412; and
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((4-trifluoromethylphenyl)-piperazin-1-yl) pteridine (example 180) was obtained from 1-(4-trifluoromethylphenyl)-piperazine in 48 % yield and characterised as follows: $R_f = 0.55$ (MeOH/CH₂Cl₂=1/9); UV (MeOH/H₂O, nm): 212, 266, 294, 412; MS (m/z): 512.2 ([M+H]⁺, 100).

Examples 181 to 184 - synthesis of 2-amino-6-(3,4-dimethoxyphenyl)-4-(substituted piperazin-1-yl) pteridine trihydrochloride salts

A mixture of the compound of example 4 (299 mg, 1.0 mmole), 1,1,1,3,3,3-hexamethyldisilazane (1 ml, 4.7 mmole), a N-substituted piperazine (4.0 mmole), p-toluenesulfonic acid (20 mg, 0.1 mmole) and ammonium sulfate (20 mg, 0.15 mmole) in toluene (4 ml) was refluxed for 48 hours (the reaction mixture became clear when the reaction was finished). After removing the solvents under reduced pressure, the residue was purified by flash chromatography over silica (CH₃OH/CH₂Cl₂ 1:20 to 1:30). To a solution of the resulting free pteridine base in methanol (20 ml), 1.25 M HCl in MeOH (4 ml, 5.0 mmole) was slowly added. The mixture was stirred at room temperature for one hour. The precipitate (which is the trihydrochloride salt of the free pteridine base) was filtered off and washed with methanol. Drying in the vacuum over P₂O₅ yielded the corresponding hydrochloride salt as a yellow solid. The following salts were made according to this procedure, with yields indicated below:

- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-dimethylaminoethyl)-piperazin-1-yl)-pteridine trihydrochloride salt (example 181) obtained in 58% yield was characterised as follows:

R_f = 0.20 (MeOH/Et₃N/CH₂Cl₂=4/2/100); UV (MeOH/H₂O, nm): 215, 297, 412; MS (m/z): 439 ([M-3HCl+H]⁺, 100);

- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(3-dimethylaminopropyl)-piperazin-1-yl) pteridine trihydrochloride salt (example 182) obtained in 57 % yield was characterised as follows:

5 R_f = 0.25 (MeOH/Et₃N/CH₂Cl₂=4/2/100); UV (MeOH/H₂O, nm): 214, 296, 412; MS (m/z): 453 ([M-3HCl+H]⁺, 100);

- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-dipropylaminoethyl)-piperazin-1-yl) pteridine trihydrochloride salt (example 183) obtained in 52 % yield was characterised as follows:

10 R_f = 0.40 (MeOH/Et₃N/CH₂Cl₂=4/2/100); UV (MeOH/H₂O, nm): 216, 297, 413; MS (m/z): 495 ([M-3HCl+H]⁺, 100); and

- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-piperidin-1-yl-ethyl)-piperazin-1-yl) pteridine trihydrochloride salt (example 184) obtained in 44 % yield was characterised as follows:

15 R_f = 0.35 (MeOH/Et₃N/CH₂Cl₂=4/2/100); UV (MeOH/H₂O, nm): 216, 297, 412; MS (m/z): 479 ([M-3HCl+H]⁺, 100).

Examples 185 to 187 - synthesis of 2-amino-4-(N-substituted-piperazino)-6-(3,4-dimethoxyphenyl) pteridines

While repeating the experimental procedure of examples 64-66 and 133-162, the three following compounds were obtained as yellow powders:

- 20 - 2-amino-4-[4-trifluoromethyl-2-nitro-phenyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 185) was obtained from 1-(4-trifluoromethyl-2-nitro-phenyl)-piperazine and characterised as follows : MS m/z (%) 557 ([M+H]⁺, 100) ;
- 2-amino-4-[2-trifluoromethyl-4-nitro-phenyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine (example 186) was obtained from 1-(2-trifluoromethyl-4-nitro-phenyl)-piperazine and characterised as follows : MS m/z (%) 557 ([M+H]⁺, 100) ; and
- 25 - 2-amino-4-[2-(piperazin-1-yl)-acetic acid N-(2-thiazolyl)-amide]-6-(3,4-dimethoxyphenyl) pteridine (example 187) was obtained from 2-(piperazin-1-yl)-acetic acid N-(2-thiazolyl)-amide and characterised as follows : MS m/z (%) 508 ([M+H]⁺, 100).

Example 188 – mixed lymphocyte reaction assay

30 Pteridine derivatives were first dissolved (10mM) in dimethylsulfoxide (hereinafter referred as DMSO) and further diluted in culture medium before use for the following *in vitro* experiments. The commercially available culture medium consisted of RPMI-1640 + 10% foetal calf serum (FCS). Some pteridine derivatives described in the previous examples (as indicated in table 1) were tested in the following mixed lymphocyte reaction (MLR) assay.

35 Peripheral blood mononuclear cells (hereinafter referred as PBMC) were isolated from heparinized peripheral blood by density gradient centrifugation over Lymphoprep (Nycomed, Maorstua, Norway). Allogeneic PBMC or Eppstein-Barr Virus-transformed human B cells [commercially available under the trade name RPMI1788 (ATCC name CCL156)]

which strongly express B7-1 and B7-2 antigens were used as stimulator cells after irradiation with 30 Gy. MLR was performed in triplicate wells. After 5 days incubation at 37°C, 1 µCi [³H]-thymidine was added to each cup. After a further 16 hours incubation, cells were harvested and counted in a β-counter. Inhibition of proliferation by a compound (drug) described in some of the previous examples was counted using the formula:

$$\% \text{ inhibition} = \frac{(\text{cpm} + \text{drugs}) - (\text{cpm cult. med})}{(\text{cpm} - \text{drugs}) - (\text{OD cult. med})} \times 100$$

wherein cpm is the thymidine count per minute. The MLR assay is regarded by those skilled in the art as an *in vitro* analogue of the transplant rejection since it is based on the recognition of allogeneic major histocompatibility antigens on the stimulator leukocytes, by responding lymphocytes.

Table 1 below shows the IC₅₀ values for various pteridine derivatives in the MLR test. The IC₅₀ value represents the lowest concentration of the pteridine derivative (expressed in µmole/l) that resulted in a 50% suppression of the MLR.

TABLE 1

Example n°	MLR	Example n°	MLR	Example n°	MLR
8	0.9	29	0.0005	49	0.4
9	0.5	30	0.1	50	0.065
10	0.1	31	3.0	51	0.3
11	0.4	32	0.4	55	5.1
12	0.3	36	0.8	56	2.8
13	0.1	37	< 0.001	57	1.9
14	0.2	38	0.3	58	7.6
15	0.1	39	0.7	60	4.1
16	0.8	40	0.6	62	10
17	0.0038	41	0.5	63	10
18	0.3	42	0.4	64	0.4
19	2.4	43	0.2	65	0.9
20	0.5	44	0.2	66	0.8
21	6.0	45	0.09	79	4.1
26	0.5	46	0.6	80	4.7
27	0.08	47	0.3	82	3.2
28	0.03	48	0.3	87	8.4
89	0.087	90	0.058	91	0.052
92	0.05	93	0.07	94	0.1
95	0.12	96	0.15	97	0.24

TABLE 1 (follow)

Example n°	MLR	Example n°	MLR	Example n°	MLR
98	0.12	99	0.0034	100	0.0074
101	0.0008	102	0.074	103	0.0065
104	0.018	105	0.037	106	0.058
107	0.003	108	0.034	109	0.1
111	8.8	118	0.9	119	0.3
122	0.8	123	3.9	124	0.9
125	0.8	126	4.9	127	9.1
128	7.4	129	2.5	130	1.5
131	0.3	132	< 0.1	133	1.2
136	1.4	137	7.6	138	6.0
141	0.7	144	4.3	145	3.5
146	0.2	147	0.2	148	0.3
149	0.5	150	1.0	151	0.6
152	1.2	153	0.2	154	0.1
155	3.6	156	0.6	157	0.5
158	0.8	159	0.11	160	3.8
161	0.4	162	0.8	135	0.6
163	0.5	164	2.3	165	2.9
166	5.3	167	3.8	168	1.6
169	0.8	171	3.4	172	0.1
173	0.7	174	0.5	175	0.4
176	0.3	177	0.8	178	1.3
179	7.9	183	8.2	184	7.7
185	1.9	186	5.0	187	0.1

5 Example 189 - TNF- α assay

Peripheral blood mononuclear cells (herein referred as PBMC), in response to stimulation by lipopolysaccharide (hereinafter LPS), a gram-negative bacterial endotoxin, produce various chemokines, in particular human TNF- α . Inhibition of the activation of PBMC can therefore be measured by the level of suppression of the production of TNF- α by PBMC in response to stimulation by LPS.

Such inhibition measurement was performed as follows: PBMC were isolated from heparinized peripheral blood by density gradient centrifugation over Lymphoprep (commercially available from Nycomed, Norway). LPS was then added to the PMBC suspension in complete medium (10^6 cells /ml) at a final concentration of 1 μ g/ml. The pteridine derivative to be tested was added at different concentrations (0.1 μ M, 1 μ M and 10 μ M) and the cells were incubated at 37°C for 72 hours in 5% CO₂. The supernatants were collected, then TNF- α concentrations were measured with respectively an anti-TNF- α antibody in a sandwich ELISA (Duo Set ELISA human TNF α , commercially available from R&D Systems, United Kingdom). The colorimetric reading of the ELISA was measured by a Multiskan RC plate reader (commercially available from ThermoLabsystems, Finland) at 450 nm (reference wavelength: 690 nm). Data analysis was performed with Ascent software 2.6. (also from ThermoLabsystems, Finland): a standard curve (recombinant human TNF α) was drawn and the amount (pg/ml) of each sample on the standard curve was determined.

The % suppression of human TNF α production by the pteridine derivatives of the invention (drugs) was calculated using the formula:

$$\% \text{ suppression} = \frac{\text{pg/ml in drugs} - \text{pg/ml in cult. med.}}{(\text{pg/ml in cult. med.} + \text{LPS}) - \text{pg/ml cult. med.}}$$

Table 2 below shows the IC₅₀ values (expressed in μ M) of the tested pteridine derivatives in the TNF- α assay.

TABLE 2

Example n°	TNF- α	Example n°	TNF- α	Example n°	TNF- α
8	0.4	29	0.077	48	0.08
9	0.14	30	0.3	49	0.07
10	0.07	31	0.5	50	0.02
11	0.4	32	0.4	51	0.04
12	0.2	36	0.3	55	10
13	0.08	37	0.1	56	3.7
14	0.06	38	0.06	60	2.5
15	0.01	39	0.8	62	7.6
16	0.55	40	0.7	64	0.05
17	0.08	41	0.9	65	0.095
18	0.09	42	0.1	66	0.3
19	0.06	43	0.7	79	9.8
20	0.03	44	0.7	80	10.0

TABLE 2 (Follow)

Example n°	TNF- α	Example n°	TNF- α	Example n°	TNF- α
26	0.5	45	0.4	82	9.3
27	0.5	46	0.2	89	1.2
28	0.5	47	0.6	90	0.04
91	0.06	92	0.06	93	0.05
94	0.09	95	0.1	96	0.01
97	0.3	98	0.08	99	0.09
100	0.51	101	0.43	102	0.63
103	0.1	104	0.6	105	0.7
106	1.4	107	0.5	108	0.4
109	0.4	112	0.28	113	3.2
119	0.26	122	0.6	123	0.8
124	0.4	126	0.9	127	2.6
128	3.3	129	0.8	130	1.4
131	1.9	132	0.05	133	0.33
134	0.07	135	0.27	136	0.8
		137	0.7	139	0.4
140	1.3	141	0.2	143	0.8
144	0.7	145	4.4	146	0.2
147	0.1	148	0.3	149	0.3
150	0.3	151	0.1	152	0.3
153	0.1	154	0.03	155	0.9
156	0.4	157	0.3	158	0.7
159	0.02	160	0.42	161	0.06
162	0.4	163	0.5	164	0.2
165	0.3	166	0.8	167	0.3
168	0.09	169	0.3	170	0.4
171	0.5	172	0.05	174	0.11
176	< 0.01	177	0.09	178	0.15
179	1.8	180	1.4	181	8.8
182	4.8	183	1.2	184	0.7
185	0.5	186	0.4	187	0.06

Example 190 - protection against a lethal dose of TNF- α

A model of TNF- α induced shock in C57BL/6 male mice was performed as follows. Five animals from the control group received an intravenous administration of a lethal dose of TNF- α (10 μ g) in the tail. Ten animals from the test group received three intraperitoneous injections of the pteridine derivative of example 17 (20 mg/kg/day) respectively 48 hours, 24 hours and immediately before an intravenous injection of TNF- α (10 μ g).

Body temperature, a clinical sign of TNF-induced shock, was followed for 40 hours in control mice and in mice receiving the pteridine derivative of example 17: the body temperature of control mice dropped significantly when compared to mice receiving the test compound of example 17.

Furthermore, all five mice from the control group died within 40 hours, the survival rate (80%) of mice that received the pteridine derivative of example 17 in addition to the TNF- α dose (10 μ g) was quite substantial.

Example 191 – inhibition of the metastasis of melanoma cells in mice

C57BL/6 mice were injected with 1.5×10^6 B16BL/6 melanomasarcoma cells subcutaneously in the foot path and were divided, three days later, into 4 groups:

- a control 1 group of 6 mice receiving only vehicle 3 times a week;
- a control 2 group of 5 mice receiving a lethal dose of TNF (15 μ g, subcutaneously) 3 times a week;
- a group 3 receiving the pteridine derivative of example 17 alone 3 times a week at 20mg/kg
- a group 4 of 7 mice receiving the pteridine derivative of example 17 in an amount of 20 mg/kg intraperitenously on days 3, 4, and 5 and also receiving TNF (15 μ g, subcutaneously) on day 5. This combined treatment with TNF and the pteridine derivative of example 17 was continued for 2 weeks.

Tumor size data (tumor size was measured as the largest diameter multiplied by the smallest diameter) show that the combined treatment in group 4 leads to a significant reduction of tumor size (120 mm²) when compared to the control group 1 (tumor size: 440 mm²). Reduction of tumor size is also true in mice of group 3, although to a lesser extent (198 mm²).

All mice of control group 2 died within the very first days of treatment, whereas mortality was 3/7 in mice of group 4.

At the end of experiment, it was looked macroscopically at black metastasis in inguinal and/or para-aortic lymphoneuds in all tumor bearing groups of mice. The proportions of mice having metastasis were:

- 4/5 mice in control group 1,
- 0/4 mice in group 4, and
- 2/4 in group 3.

Example 192 - IL-1 β assay

Peripheral blood mononuclear cells (herein referred as PBMC), in response to stimulation by lipopolysaccharide (LPS), a gram-negative bacterial endotoxin, produce various chemokines, in particular human IL-1 β . Inhibition of the activation of PBMC can therefore be measured by the level of suppression of the production of IL-1 β by PBMC in response to stimulation by LPS.

Such inhibition measurement was performed as follows: PBMC were isolated from heparinized peripheral blood by density gradient centrifugation over Lymphoprep (commercially available from Nycomed, Norway). LPS was then added to the PMBC suspension in complete medium (10^6 cells /ml) at a final concentration of 1 μ g/ml. The pteridine derivative to be tested was added at different concentrations (0.1 μ M, 1 μ M and 10 μ M) and the cells were incubated at 37°C for 72 hours in 5% CO₂. The supernatants were collected, then IL-1 β concentrations were measured with an anti-IL-1 β antibody in a sandwich ELISA. The colorimetric reading of the ELISA was measured by a Multiskan RC plate reader (commercially available from ThermoLabsystems, Finland) at 450 nm (reference wavelength: 690 nm). Data analysis was performed with Ascent software 2.6. (also from ThermoLabsystems, Finland) : a standard curve (recombinant human IL-1 β) was drawn and the amount (pg/ml) of each sample on the standard curve was determined.

The % suppression of human IL-1 β by the pteridine derivatives (drugs) of this invention was calculated using the formula:

$$\% \text{ suppression} = \frac{\text{pg/ml in drugs} - \text{pg/ml in cult. med.}}{(\text{pg/ml in cult. med.} + \text{LPS}) - \text{pg/ml cult. med.}}$$

Table 3 below shows the IC₅₀ values (expressed in μ M) of the tested pteridine derivatives in the IL-1 β assay.

TABLE 3

Example n°	IL-1 β	Example n°	IL-1 β	Example n°	IL-1 β
12	5.0	15	7.0	17	2.8
20	8.5	21	6.7	36	0.8
37	3.3	42	0.9	46	0.8
47	0.9	48	0.6	49	2.2
50	5.5	56	8.7	57	9.5
64	4.4	65	3.5	91	4.2
92	8.6	93	4.5	95	1.0
98	6.2	100	7.4	101	5.6
107	0.8	108	1.0	122	10
123	4.2	124	5.0	125	7.5

126	3.7	132	5.3	134	1.2
136	4.7	139	1.2	141	0.6
146	5.0	147	4.3	148	7.2
149	4.9	150	3.5	151	3.7
152	3.6	153	2.0	154	2.2
157	2.3	159	5.4	161	3.2
162	3.8	166	4.4	167	3.7
172	5.1	173	4.6	174	5.0
176	4.9	178	4.0	180	4.4
187	2.4				

Table 3 (end)

5

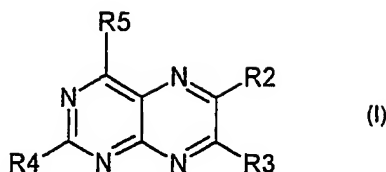
10

15

20

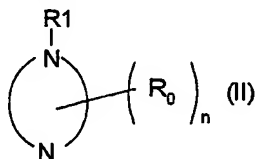
CLAIMS

1. A pteridine derivative having the general formula (I):

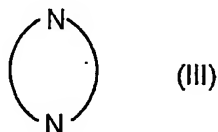


wherein:

- a first group of one or more of the substituents R_2 , R_3 , R_4 and R_5 of the pteridine ring is independently selected from groups represented by the general formula (II):



wherein:



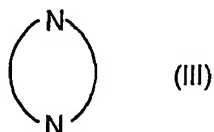
- schematically represents a saturated or partly unsaturated heterocyclic ring with at least two nitrogen atoms in the said heterocyclic ring and with a total of 5 to 7 atoms in the said heterocyclic ring, and optionally with one or more other heteroatoms (e.g. oxygen or sulfur) in the said heterocyclic ring or attached to one or more carbon atoms of said heterocyclic ring (for instance in the form of a carbonyl or thiocarbonyl group), wherein one of said at least two nitrogen atoms in the heterocyclic ring is attached to a carbon atom of the pteridine ring at any of positions 2, 4, 6 or 7 of the pteridine ring, wherein the said heterocyclic ring may be fused to one or more aromatic hydrocarbon rings, and wherein:
- 15 - each substituent R_0 of the heterocyclic ring (III) is a group independently selected from the group consisting of halogen, nitro, C_{1-7} alkyl (optionally containing one or more functions or radicals selected from the group consisting of halogen, carbonyl, thiocarbonyl, hydroxyl, sulfhydryl, C_{1-7} alkoxy, thio C_{1-7} alkyl, thio C_{3-10} cycloalkyl, acetal, thioacetal, imino, oximino, alkyloximino, amino-acid, cyano, (thio)carboxylic acid,
- 20 (thio)carboxylic acid ester or amide, nitro, amino, C_{1-7} alkylamino, cycloalkylamino, alkenylamino, cycloalkenyl-amino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercapto-alkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl and sulfonamido), C_{3-7} alkenyl, C_{2-7} alkynyl, halo C_{1-7} alkyl, C_{3-10}
- 25 cycloalkyl, aryl, arylalkyl, alkylaryl, alkylacyl, arylacyl, hydroxyl, sulfhydryl, amino, C_{1-7}

- alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkyl-amino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, C₁₋₇ alkoxy, C₃₋₁₀ cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, thioaryl, thioheterocyclic, arylalkylthio, heterocyclic-substituted alkylthio, formyl, hydroxylamino, cyano, (thio)carboxylic acid or esters or thioesters or amides or thioamides thereof;
- n is an integer from 0 to 6;
 - R₁ is a substituent group selected from the group consisting of formyl, acyl, thio-acyl, amide, thioamide, sulfonyl, sulfinyl, carboxylate, thiocarboxylate, and amino-substituted acyl, alkoxyalkyl, C₃₋₁₀ cycloalkyl-alkyl, C₃₋₁₀ cyclo-alkyl, dialkylaminoalkyl, heterocyclic-substituted alkyl, acyl-substituted alkyl, thioacyl-substituted alkyl, amido-substituted alkyl, thioamido-substituted alkyl, carboxylate-substituted alkyl, thiocarboxylate-substituted alkyl, (amino-substituted acyl)alkyl, heterocyclic, carboxylic acid ester, ω-cyanoalkyl, ω-carboxylic ester-alkyl, halo C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, arylalkenyl, aryloxyalkyl, arylalkyl and aryl, wherein the aryl moiety of each of said arylalkenyl, aryloxyalkyl, arylalkyl and aryl radicals is optionally substituted with one or more substituents independently selected from the group consisting of halogen, C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, halo C₁₋₇ alkyl, nitro, hydroxyl, sulfhydryl, amino, C₁₋₇ alkoxy, C₃₋₁₀ cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, thioaryl, thio-heterocyclic, arylalkylthio, heterocyclic-substituted alkylthio, formyl, carbamoyl, thiocarbamoyl, ureido, thioureido, sulfon-amido, hydroxylamino, alkoxyamino, mercaptoamino, thioalkyl-amino, acylamino, thioacylamino, cyano, carboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, thiocarboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, alkylamino, cycloalkylamino, alkenyl-amino, cycloalkenylamino, alkynylamino, arylamino, aryl-alkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic amino, hydrazino, alkylhydrazino and phenylhydrazino; and
 - wherein the remaining of, i.e. a second group of, the substituents R₂, R₃, R₄ and R₅ of the pteridine ring is independently selected from the group consisting of hydrogen; halogen; C₁₋₇ alkyl; C₂₋₇ alkenyl; C₂₋₇ alkynyl; halo C₁₋₇ alkyl; carboxy C₁₋₇ alkyl; carboxyaryl; C₁₋₇ alkoxy; C₃₋₁₀ cycloalkoxy; aryloxy; arylalkyloxy; oxyheterocyclic; heterocyclic-substituted alkyloxy; thio C₁₋₇ alkyl; thio C₃₋₁₀ cycloalkyl; thioaryl; thioheterocyclic; arylalkylthio; heterocyclic-substituted alkylthio; hydroxylamino; mercapto-amino; acylamino; thio-acylamino; alkoxyamino; thioalkyl-amino; acetal; thio-acetal; carboxylic acid; carboxylic acid esters, thioesters, halides, anhydrides, amides and thioamides; thiocarboxylic acid; thiocarboxylic acid esters, thioesters, halides, anhydrides, amides

and thioamides; hydroxyl; sulfhydryl; nitro; cyano; carbamoyl; thiocarbamoyl; ureido; thioureido; amino; alkylamino; cycloalkylamino; alkenylamino; cyclo-alkenylamino; alkynylamino; arylamino; arylalkylamino; hydroxyalkylamino; mercaptoalkyl-amino; heterocyclic amino; heterocyclic-substituted arylamino; heterocyclic-substituted alkyl-amino; oximino; alkyloximino; hydrazino; 5 alkylhydrazino; phenylhydrazino; cysteinyl acid, esters, thioesters, halides, anhydrides, amides and thioamides thereof; aryl optionally substituted with one or more substituents selected from the group consisting of halogen, C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, halo C₁₋₇ alkyl, nitro, hydroxyl, sulfhydryl, amino, C₁₋₇ alkoxy, C₃₋₁₀ cycloalkoxy, aryloxy, arylalkyloxy, oxyheterocyclic, heterocyclic-substituted alkyloxy, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, thioaryl, thioheterocyclic, 10 arylalkylthio, heterocyclic-substituted alkylthio, formyl, carbamoyl, thiocarbamoyl, ureido, thioureido, sulfonamido, hydroxylamino, alkoxyamino, mercaptoamino, thioalkylamino, acylamino, thioacylamino, cyano, carboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, thiocarboxylic acid or esters or thioesters or halides or anhydrides or amides thereof, alkylamino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, 15 arylalkylamino, hydroxyalkyl-amino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, hetero-cyclic-substituted arylamino, hydrazino, alkylhydrazino and phenylhydrazino; optionally substituted heterocyclic radicals; aromatic or heterocyclic substituents substituted with an aliphatic spacer between the pteridine ring and the aromatic or heterocyclic substituent, whereby said aliphatic spacer is a branched or straight, saturated or unsaturated 20 aliphatic chain of 1 to 4 carbon atoms which may contain one or more functions, atoms or radicals independently selected from the group consisting of carbonyl, thiocarbonyl, hydroxyl, thiol, ether, thioether, acetal, thioacetal, amino, imino, oximino, alkyloximino, amino-acid, cyano, acylamino, thioacylamino, carbamoyl, thiocarbamoyl, ureido, thioureido, carboxylic acid or ester or thioester or halide or anhydride or amide, thiocarboxylic acid or ester or thioester or halide or anhydride or 25 amide, nitro, thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, hydroxylamino, mercaptoamino, alkylamino, cycloalkylamino, alkenylamino, cycloalkenyl-amino, alkynylamino, arylamino, arylalkylamino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, heterocyclic amino, heterocyclic-substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl, sulfinyl, sulfonamido and halogen; branched or straight, saturated or unsaturated aliphatic chains of 1 to 7 30 carbon atoms optionally containing one or more functions, atoms or radicals independently selected from the group consisting of halogen, carbonyl, thiocarbonyl, hydroxyl, thiol, ether, thioether, acetal, thio-acetal, amino, imino, oximino, alkyloximino, aminoacid, cyano, acylamino; thioacylamino; carbamoyl, thiocarbamoyl, ureido, thio-ureido, carboxylic acid ester or halide or anhydride or amide, thiocarboxylic acid or ester or thioester or halide or anhydride or amide, nitro, 35 thio C₁₋₇ alkyl, thio C₃₋₁₀ cycloalkyl, hydroxylamino, mercaptoamino, alkyl-amino, cycloalkylamino, alkenylamino, cycloalkenylamino, alkynylamino, arylamino, arylalkyl-amino, hydroxyalkylamino, mercaptoalkylamino, heterocyclic-substituted alkylamino, hetero-cyclic amino, heterocyclic-

- substituted arylamino, hydrazino, alkylhydrazino, phenylhydrazino, sulfonyl, sulfinyl and sulfonamido; or R_2 together with R_3 and the carbon atoms in positions 6 and 7 of the pteridine ring forms a homocyclic or heterocyclic radical; with the first proviso that when R_4 stands for N-formylpiperazino then R_2 may not be hydrogen, C_{1-7} alkyl or phenyl; and with the second proviso
- 5 that when R_4 stands for N-acetylpiperazino then R_2 may not be halogen; and/or a pharmaceutically acceptable addition salt thereof and/or a stereoisomer thereof and/or a mono- or a di-*N*-oxide thereof and/or a solvate thereof and/or a dihydro- or tetrahydropteridine derivative thereof.

- 10 2. A pteridine derivative according to claim 1, wherein

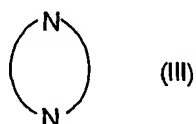


represents a piperazin-1-yl group or a homopiperazin-1-yl group, and n is 0, 1 or 2.

3. A pteridine derivative according to claim 1 or claim 2, wherein one or two substituents selected from R_4 and R_5 are groups independently having the general formula (II).

15

4. A pteridine derivative according to claim 1 or claim 2, wherein R_5 is a group having the general formula (II), wherein



represents a piperazin-1-yl group or a homopiperazin-1-yl group, and n is 0, 1 or 2.

- 20 5. A pteridine derivative according to any of claims 1 to 4, wherein R_2 is selected from the group consisting of hydrogen, C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl, preferably a phenyl group optionally substituted with one or more substituents selected from the group consisting of halogen, C_{1-7} alkyl and C_{1-7} alkoxy.
- 25 6. A pteridine derivative according to any of claims 1 to 5, wherein R_3 is selected from the group consisting of hydrogen, C_{1-7} alkyl, C_{3-10} cycloalkyl, aryl and heteroaryl.
7. A pteridine derivative according to any of claims 1 to 6, wherein R_4 is amino or a group represented by the general formula (II).

30

8. A pteridine derivative according to any of claims 1 to 7, wherein R_5 is a group selected from the group consisting of piperazin-1-yl, 2-methylpiperazin-1-yl, 2-phenylpiperazin-1-yl, homopiperazin-1-yl and 2,5-dimethylpiperazin-1-yl, the said group being substituted in its 4 position with a substituent R_1 which has a carbonyl (oxo) or thiocarbonyl (thioxo) or sulfonyl function.

5

9. A pteridine derivative according to any of claims 1 to 8, wherein R_5 is a piperazin-1-yl group substituted in its 4 position with a substituent R_1 , wherein R_1 is selected from the group consisting of:

- COR_8 wherein R_8 is selected from the group consisting of hydrogen; C_{1-7} alkyl; C_{3-10} cycloalkyl; aryl optionally substituted with one or more substituents selected from the group consisting of halogen, C_{1-7} alkyl, cyano and C_{1-7} alkoxy; heterocyclic optionally substituted with one or more halogen atoms; arylalkyl; aryloxyalkyl; arylalkoxyalkyl; alkoxyalkyl; arylalkoxy; aryloxy; arylalkenyl; heterocyclic-substituted alkyl; alkylamino and arylamino,
- CSR_9 , wherein R_9 is selected from the group consisting of alkylamino and aryloxy,
- SO_2R_{10} , wherein R_{10} is selected from the group consisting of aryl and arylalkyl, and
- R_{11} , wherein R_{11} is selected from the group consisting of C_{1-7} alkyl, aryl, arylalkyl, arylalkenyl, alkoxyalkyl, heterocyclic-substituted alkyl, cycloalkylalkyl, heterocyclic, C_{3-10} cycloalkyl, alkylaminoalkyl, aryloxyalkyl, alkoxyaryl, ω -cyanoalkyl, ω -carboxylatoalkyl and carboxamidoalkyl.

20

10. A pteridine derivative according to any of claims 1 to 9, being selected from the group consisting of:

- 2-amino-4-(*N*-acetyl piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(*N*-propionyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(*N*-hexanoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-benzoyl piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(4-chlorobenzoyl)]piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(*N*-2-thiophenecarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-[(*N*-diethylcarbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-[(*N*-hydrocinnamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(4-cyanobenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(*N*-phenoxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-[(*N*-4-butylbenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(*N*-isonicotinoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(*N*-diisopropylcarbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-[*N*-(4-pentoxybenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;

35

- 2-amino-4-[N-(3-methoxybenzoyl)piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-[N-(2-furoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(benzyloxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(p-chlorophenoxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 5 - 2-amino-4-[N-(cyclohexylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(phenylsulfonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(p-fluorobenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(2-thiophenacetyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-[N-(cinnamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 10 - 2-amino-4-[N-(1-pyrrolidinylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(diphenylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(2,6-dichloro-5-fluoro-nicotinoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(methoxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 15 - 2-amino-4-[N-(2-methoxybenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-[N-(benzylsulfonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(3,4-dichlorobenzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(4-chlorophenylacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(1-naphtoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 20 - 2-amino-4-[N-(3-furoylcarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(benzyloxycarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(dimethylthiocarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(phenoxyacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(phenoxythiocarbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 25 - 2-amino-4-[N-(2-(S)-amino-3-phenylpropionyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(2-(S)-amino-3-(4-hydroxyphenyl)propionyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(pyrrolidin-2-(S)-yl)carbonyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 30 - 2-amino-4-[N-(2-(S)-amino-3-(indol-2-yl)propionyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(phenoxyacetyl)-homopiperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[N-(4-methyl-phenoxy-carbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 35 - 2-amino-4-[N-(4-methoxy-phenoxy-carbonyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;

- 2-amino-4-[*N*-(2-methoxy)-phenoxy-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(4-chloro)-phenoxy-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 5 - 2-amino-4-[*N*-isobutoxy-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(2-chloro)-phenoxy-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(2-methoxy)-ethoxy-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 10 - 2-amino-4-[*N*-(2-naphthoxy)-carbonyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-phenyl-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(4-fluorophenyl)-carbamoyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-(4-methylphenyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 15 - 2-amino-4-(*N*-(4-cyanophenyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-(3-methylphenyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-(benzyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-(4-fluorobenzyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-(3-chloro-4-fluorophenyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 20 - 2-amino-4-(*N*-(3-thienyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(2-(2-thienyl)ethyl)-carbamoyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(*N*-butyl)-carbamoyl]-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 25 - 2-amino-4-[*N*-(aminoacetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(2-(S),4-diamino-4-oxobutanoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(4-(methoxycarbonyl)benzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 30 - 2-amino-4-[*N*-(4-(dimethylamino)acetyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(4-amino-4-oxo-butanoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(3-carboxypropanoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[*N*-(4-(carboxy)benzoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(*N*-(methyl-*N*-phenyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 35 - 2-amino-4-(*N*-(ethyl-*N*-phenyl)-carbamoyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;

- 2-amino-4-(*N*-methyl-*N*-tolyl-carbamoyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-(2-methoxyethyl)piperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-cyclohexylmethyl)piperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-cyclopentylpiperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 5 - 2-amino-4-(1-butylpiperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-isopropylpiperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-(2-diethylaminoethyl)-piperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-(2-diisopropylaminoethyl)-piperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-(2-morpholino-4-yl-ethyl)-piperazino)-6-(3,4-dimethoxyphenyl) pteridine;
- 10 - 2-amino-4-(4-[2-(piperazin-1-yl)-acetyl]-morpholino)-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-(4-[2-(piperazin-1-yl)-acetyl]-pyrrolidino)-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-(2-[piperazin-1-yl]-acetic acid *N*-methyl *N*-phenyl amide)-6-(3,4-dimethoxy-phenyl) pteridine;
- 2-amino-4-(2-(piperazin-1-yl)-propionic acid ethyl ester)-6-(3,4-dimethoxyphenyl) pteridine;
- 15 - 2-amino-4-(3-(piperazin-1-yl)-propionic acid ethyl ester)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(2-(piperazin-1-yl)-acetic acid ethyl ester)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-(3-methyl-benzyl)piperazinyl)-6-(3,4-dimethoxyphenyl) pteridine;
- 20 - 2-amino-4-[(2,6-dichlorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(4-fluorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(4-chlorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(4-methylbenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(2-fluorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 25 - 2-amino-4-[(3,4-dichlorobenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(piperonyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(4-*tert*-butylbenzyl)piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(4-pyridyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(2-pyridyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 30 - 2-amino-4-[(2-pyrimidinyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(3-methoxyphenyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-(1-(3-phenylpropyl-piperazine)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(3,4-dichlorophenyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(3-dichlorophenyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 35 - 2-amino-4-[(1-phenylethyl)-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(2-(1-pyrrolyl)-ethyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-[(2-phenoxyethyl-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;

- 2-amino-4-(1-(2-imidazol-1-yl-ethyl-piperazine)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-((3-pyridyl)-methyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-((4-pyridyl)-methyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-((1-naphthylmethyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 5 - 2-amino-4-(*N*-phenethylpiperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-((2-methoxyphenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-((4-methoxyphenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-4-((4-chlorophenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(3-propionitril)-piperazin-1-yl) pteridine;
- 10 - 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-(1,3)-dioxolan-2-yl-ethyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-ethoxyethyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(pent-3-yl-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(1-pentyl-piperazin-1-yl) pteridine;
- 15 - 2-amino-6-(3,4-dimethoxyphenyl)-4-(1-isobutyl-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((tetrahydrofurfuryl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(1,3-dioxolan-2-yl-methylpiperazin-1-yl) pteridine;
- 2-amino-4-((3,5-dichlorophenyl)-piperazin-1-yl)-6-(3,4-dimethoxyphenyl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((4-fluorophenyl)-piperazin-1-yl) pteridine;
- 20 - 2-amino-6-(3,4-dimethoxyphenyl)-4-((3-trifluoromethylphenyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((3,4-dimethylphenyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((3-methylphenyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((4-methylphenyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((2-pyridyl)methyl-piperazin-1-yl) pteridine;
- 25 - 2-amino-6-(3,4-dimethoxyphenyl)-4-(thiazol-2-yl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(1-(1-methyl-piperidin-3-yl-methyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-((4-trifluoromethylphenyl)-piperazin-1-yl) pteridine;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-dimethylaminoethyl)-piperazin-1-yl)-pteridine trihydrochloride salt;
- 30 - 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(3-dimethylaminopropyl)-piperazin-1-yl) pteridine trihydrochloride salt;
- 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-dipropylaminoethyl)-piperazin-1-yl) pteridine trihydrochloride salt;
- 35 - 2-amino-6-(3,4-dimethoxyphenyl)-4-(4-(2-piperidin-1-yl-ethyl)-piperazin-1-yl) pteridine trihydrochloride salt;

- 2-amino-4-[2-trifluoromethyl-4-nitro-phenyl-piperazin-1-yl]-6-(3,4-dimethoxyphenyl) pteridine; and
- 2-amino-4-[2-(piperazin-1-yl)-acetic acid N-(2-thiazolyl)-amide]-6-(3,4-dimethoxyphenyl) pteridine.

5

11. A pharmaceutical composition comprising as an active principle at least one pteridine derivative according to any of claims 1 to 10.

12. A pharmaceutical composition according to claim 11, further comprising one or more biologically-active drugs selected from the group consisting of immunosuppressant and/or immunomodulator drugs, antineoplastic drugs, and antiviral agents.

13. A pharmaceutical composition according to claim 11, further comprising one or more immunosuppressant drugs selected from the group consisting of cyclosporin A; substituted xanthines; pentoxifylline; daltroban, sirolimus, tacrolimus; rapamycin and derivatives thereof; leflunomide or an active metabolite or an analog thereof; mycophenolic acid and salts thereof; adrenocortical steroids; azathioprine, brequinar, gusperimus; 6-mercaptopurine; mizoribine; chloroquine; hydroxychloroquine; monoclonal antibodies with immunosuppressive properties; etanercept; infliximab; and kineret.

15

14. A pharmaceutical composition according to claim 11, further comprising one or more immunomodulator drugs selected from the group consisting of acemannan, amiprilose, buccillamine, ditiocarb sodium, imiquimod, inosine pranobex, interferon- β , interferon- γ , lentinan, levamisole, pidotimod, romurtide, platonin, procodazole, propagermanium, thymomodulin, thymopentin and ubenimex.

20

15. A pharmaceutical composition according to claim 11, further comprising one or more antineoplastic drugs selected from the group consisting of alkaloids, alkylating agents, alkyl sulfonates, aziridines, ethylenimines, methylmelamines, nitrogen mustards, nitrosoureas, antibiotics, antimetabolites, folic acid analogs, purine analogs, pyrimidine analogs, enzymes, interferon and platinum complexes.

25

16. A pharmaceutical composition according to claim 11, further comprising one or more antiviral agents selected from the group consisting of retroviral enzyme inhibitors, HIV-1 IN inhibitors, nucleoside reverse transcriptase inhibitors, zidovudine, lamivudine, didanosine, stavudine, zalcitabine, non-nucleoside reverse transcriptase inhibitors, nevirapine, delavirdine, foscarnet

30

sodium, HIV-1 protease inhibitors, saquinavir, ritonavir, indinavir, nelfinavir, acyclovir, cidofovir, cytarabine, edoxudine, famciclovir, floxuridine, ganciclovir, idoxuridine, penciclovir, sorivudine, trifluridine, valaciclovir, vidarabine, kethoxal, methisazone, moroxydine, podophyllotoxin, ribavirine, rimantadine, stallimycine, statolon, tromantadine and xenazoic acid.

5

17. A method for the prevention or treatment in a patient of a pathologic condition selected from the group consisting of:

- immune and auto-immune disorders,
- cardiovascular disorders,
- 10 - disorders of the central nervous system,
- allergic conditions,
- TNF- α related disorders,
- viral diseases, and
- cell proliferative disorders,

15 comprising the administration to the patient of an effective amount of a pharmaceutical composition comprising as an active principle at least one pteridine derivative according to claim 1 to 10.

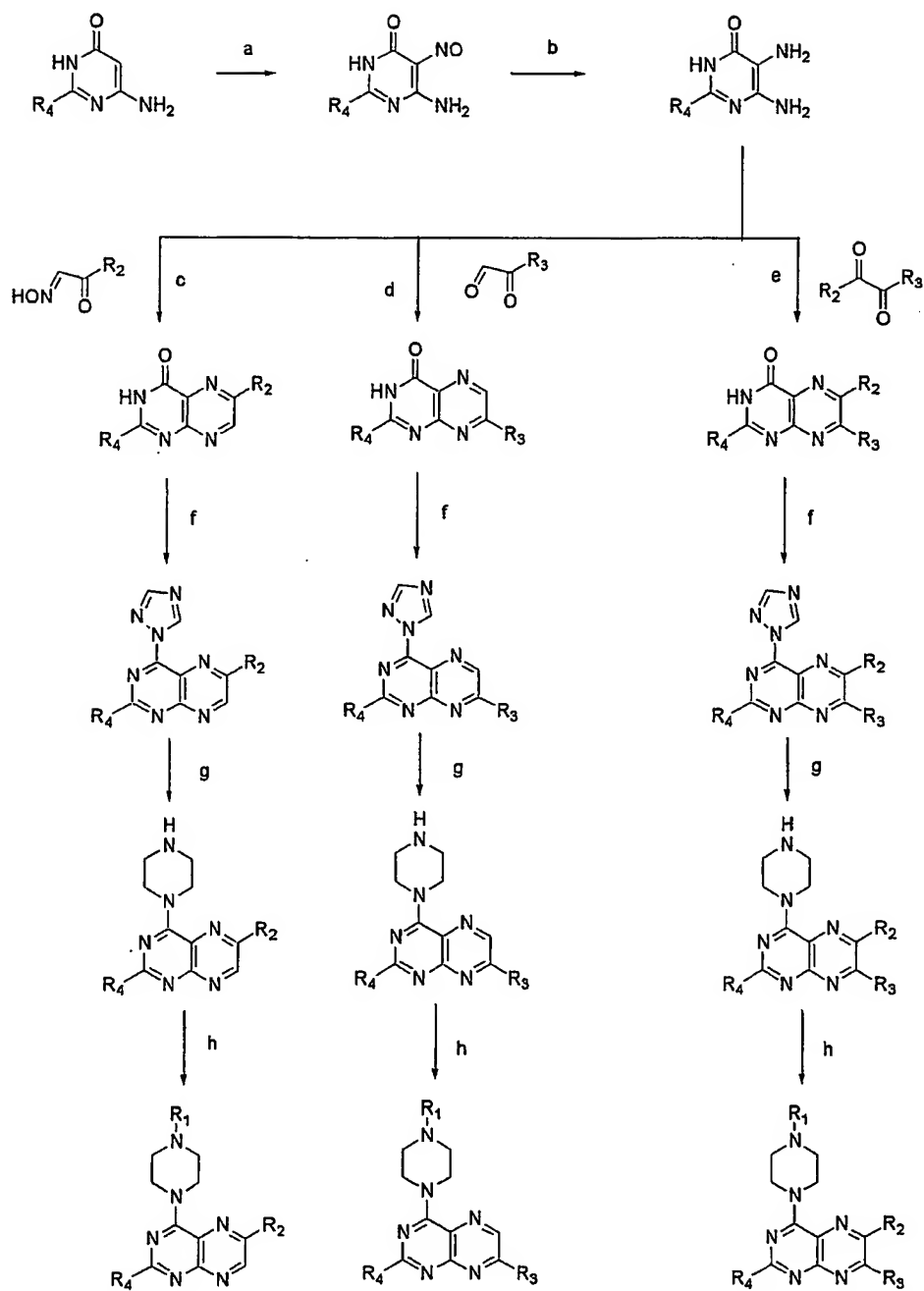
18. A method of prevention or treatment according to claim 17, wherein an effective amount of the
20 pharmaceutical composition corresponds to an amount in the range from 0.01 mg to 20 mg of the pteridine derivative per day and per kg body weight of the patient.

25

30

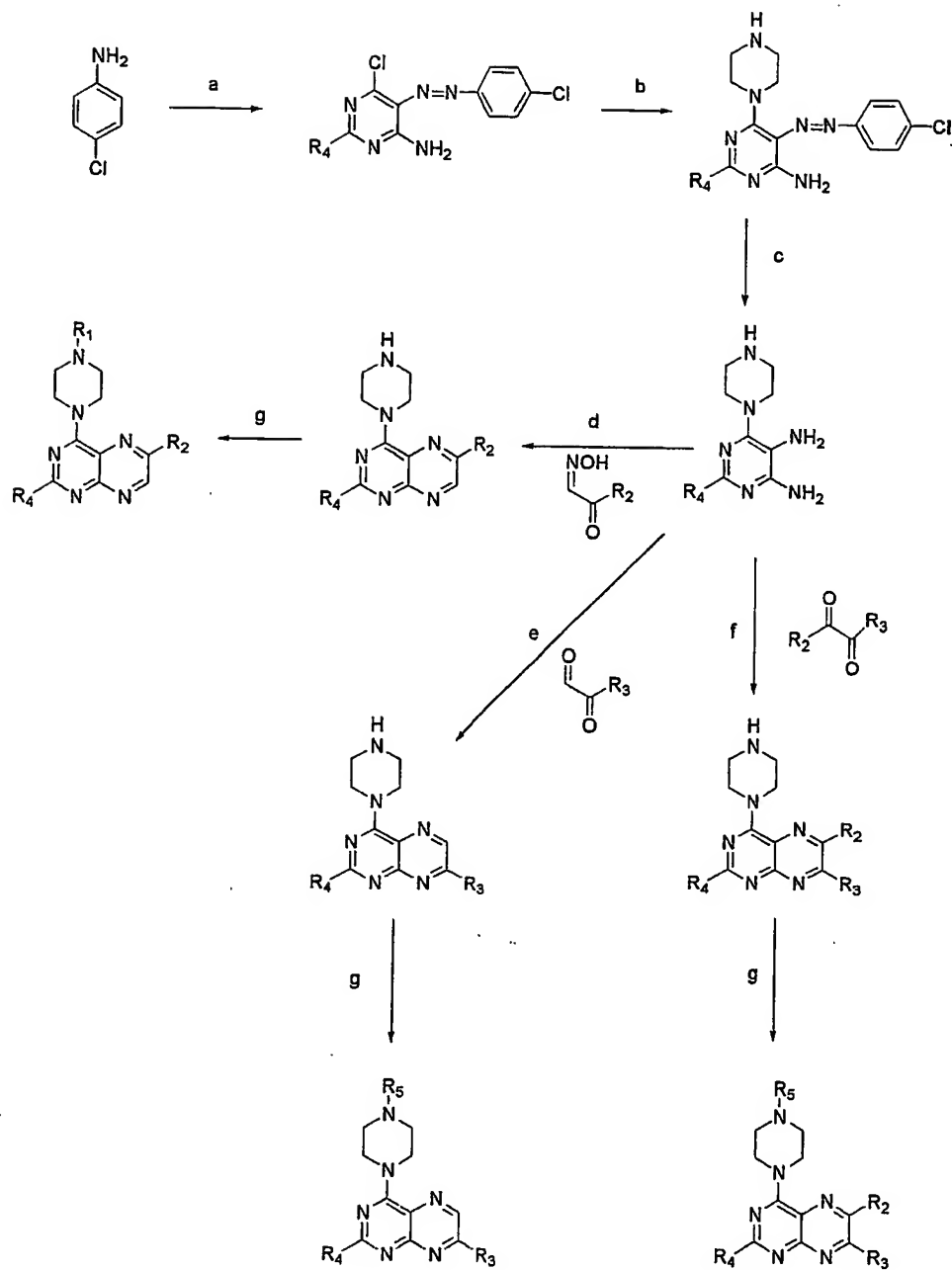
1 / 9

Figure 1



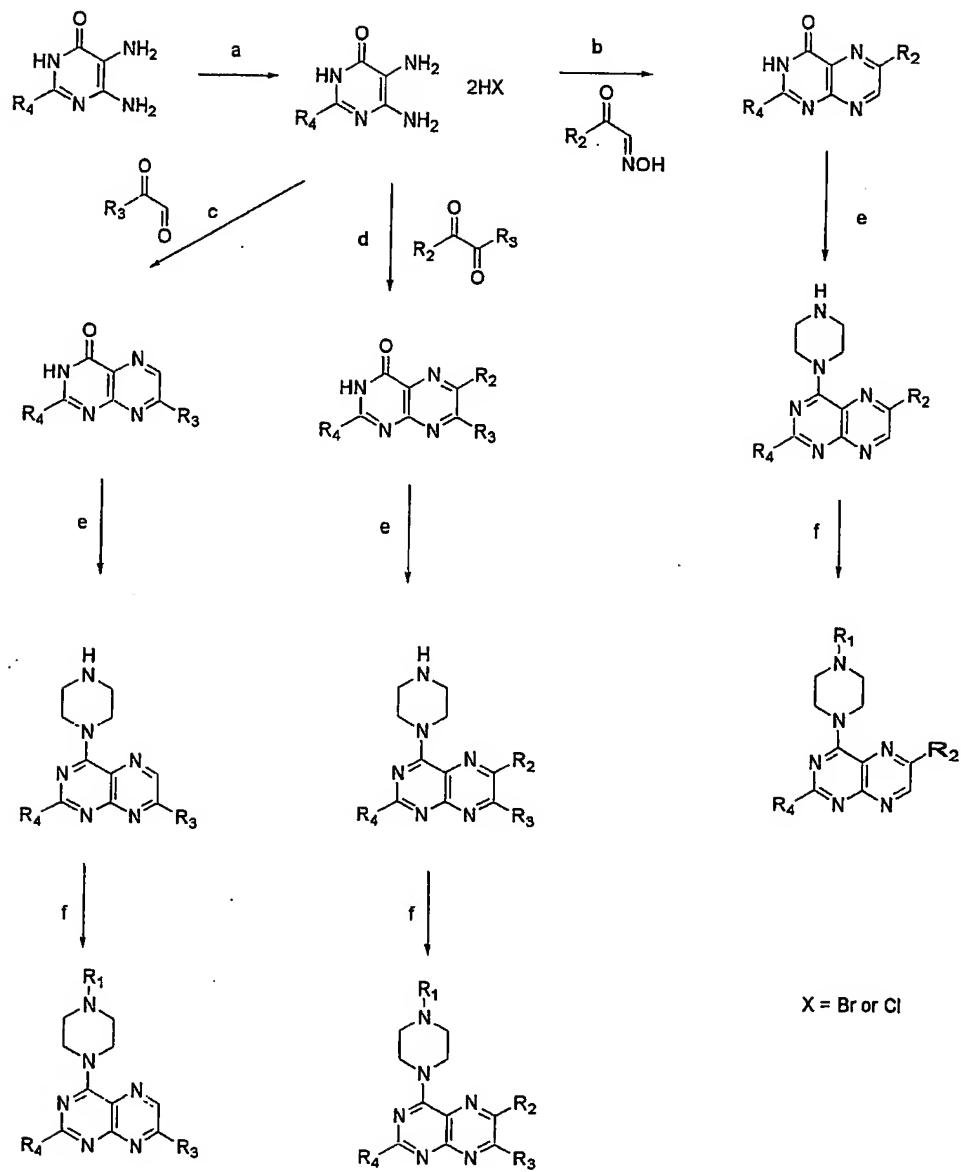
2 / 9

Figure 2



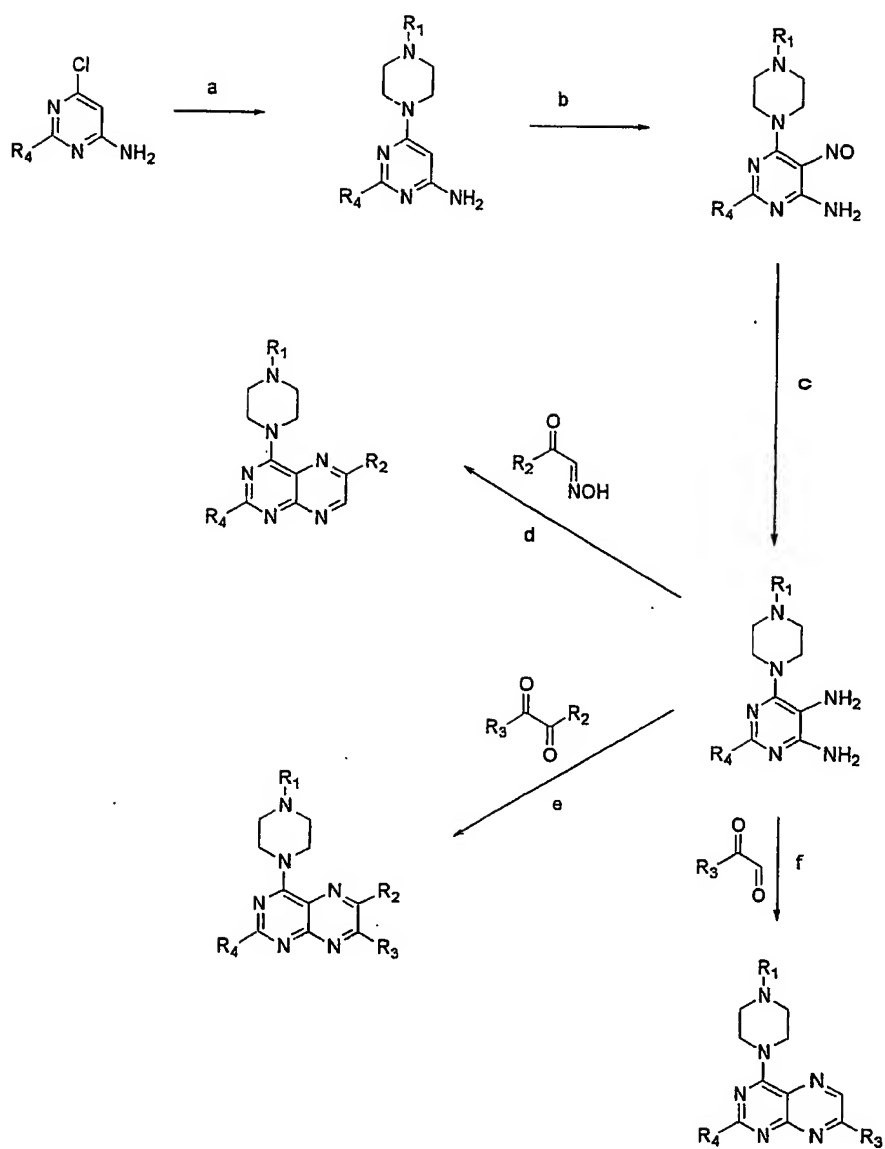
3 / 9

Figure 3



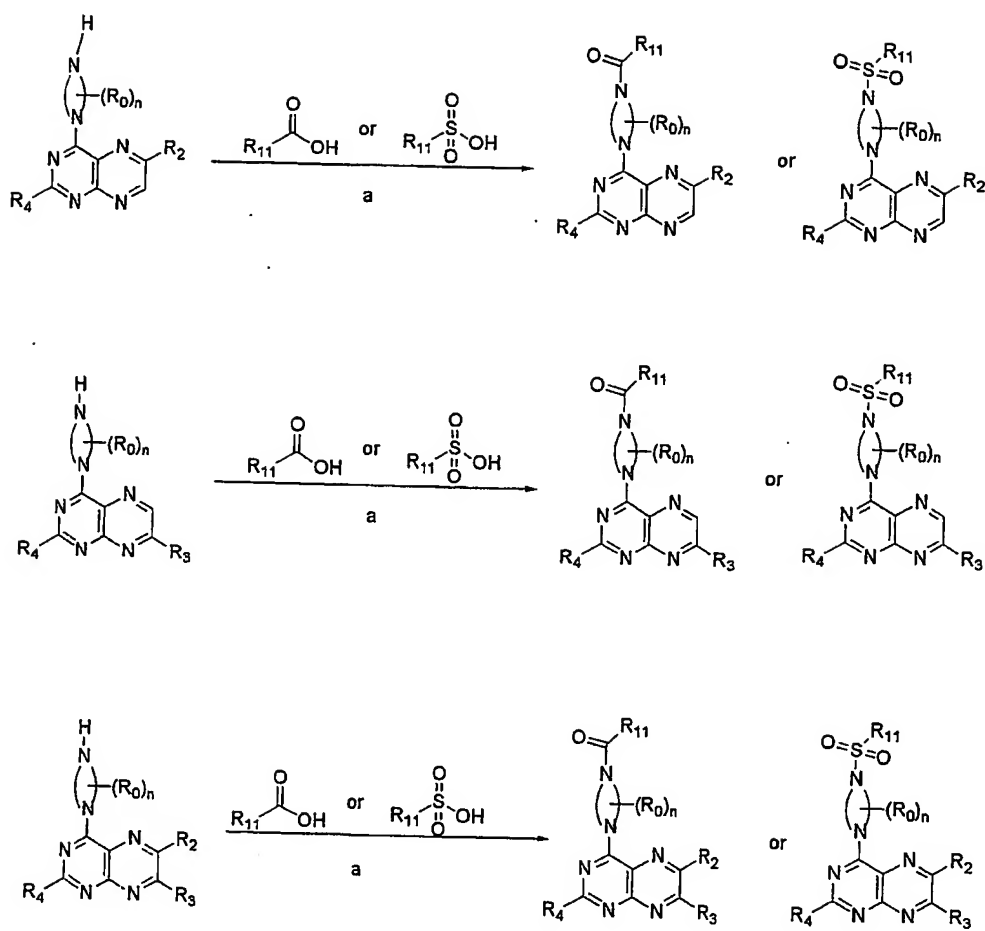
4 / 9

Figure 4



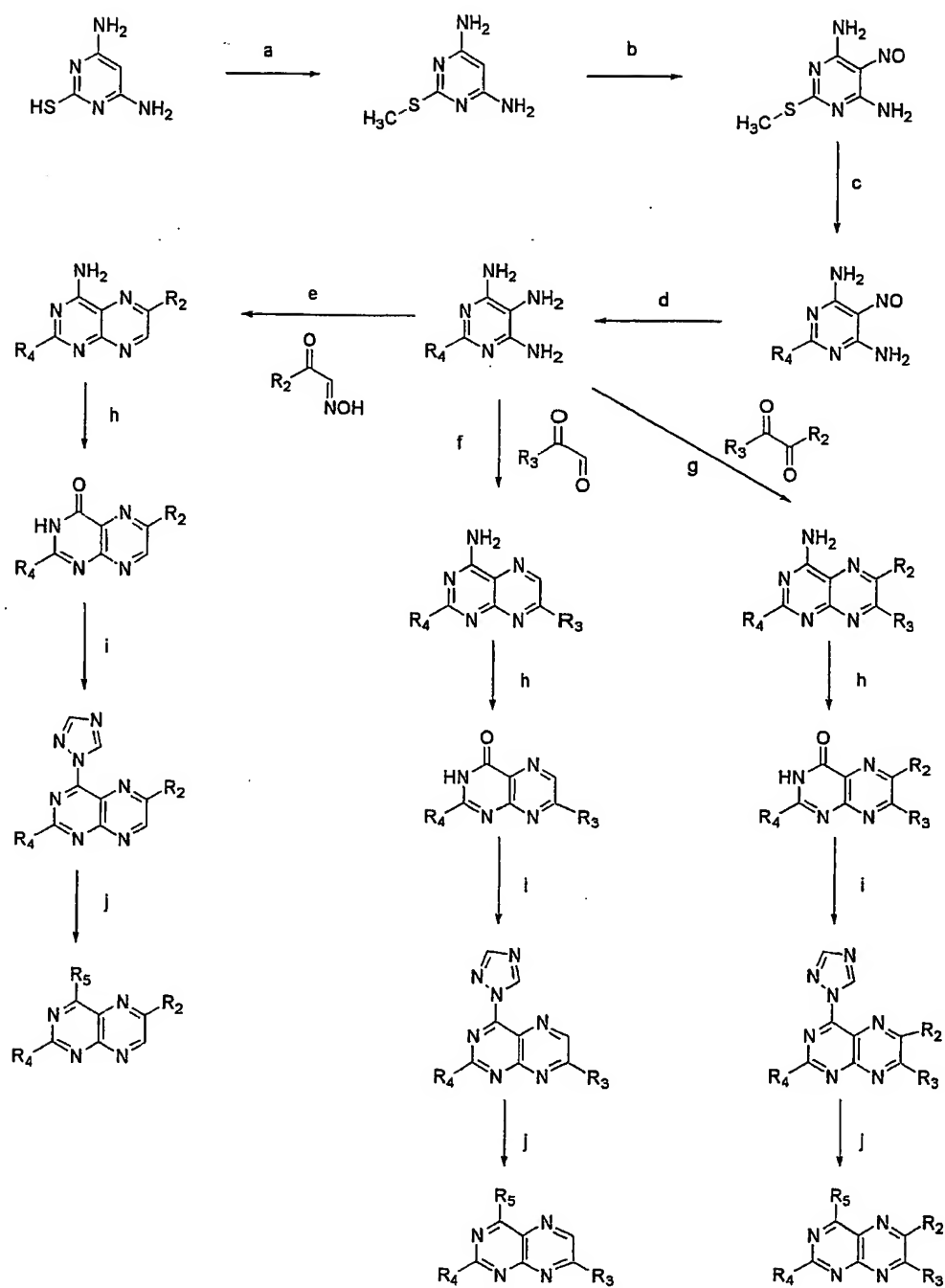
5 / 9

Figure 5



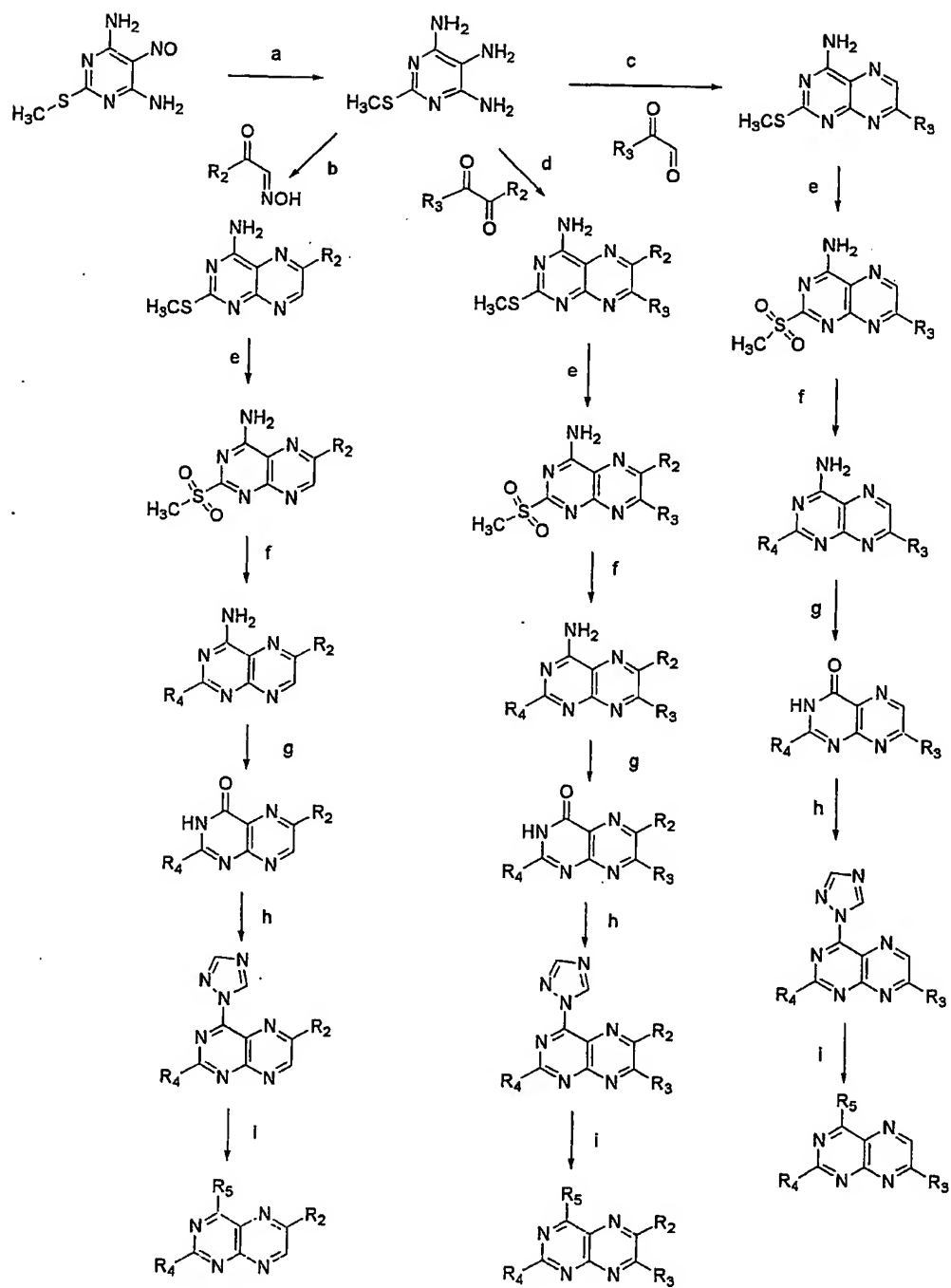
6 / 9

Figure 6



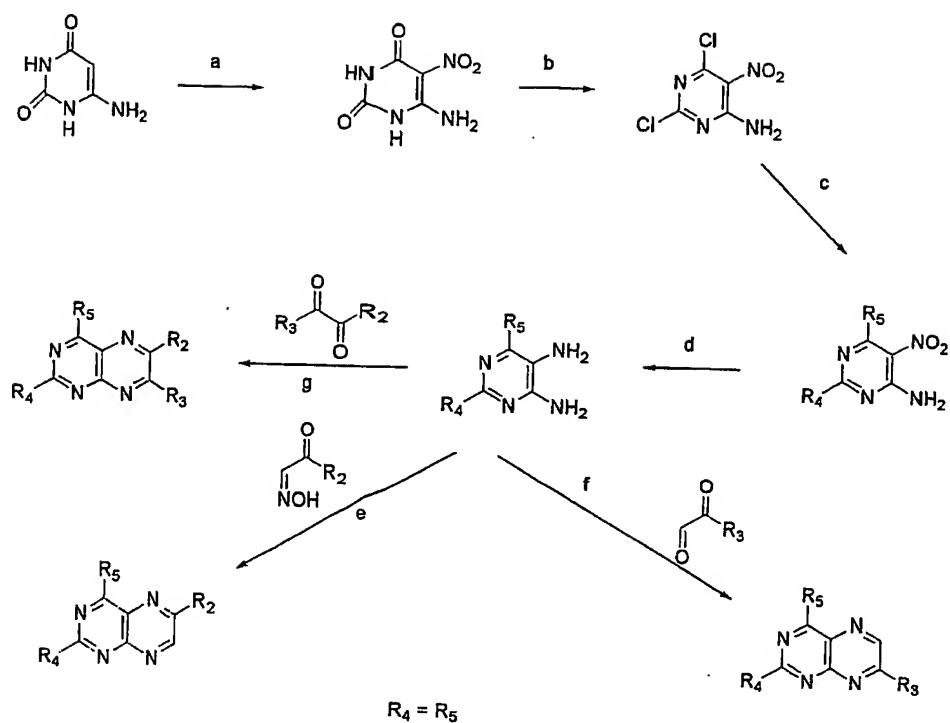
7 / 9

Figure 7



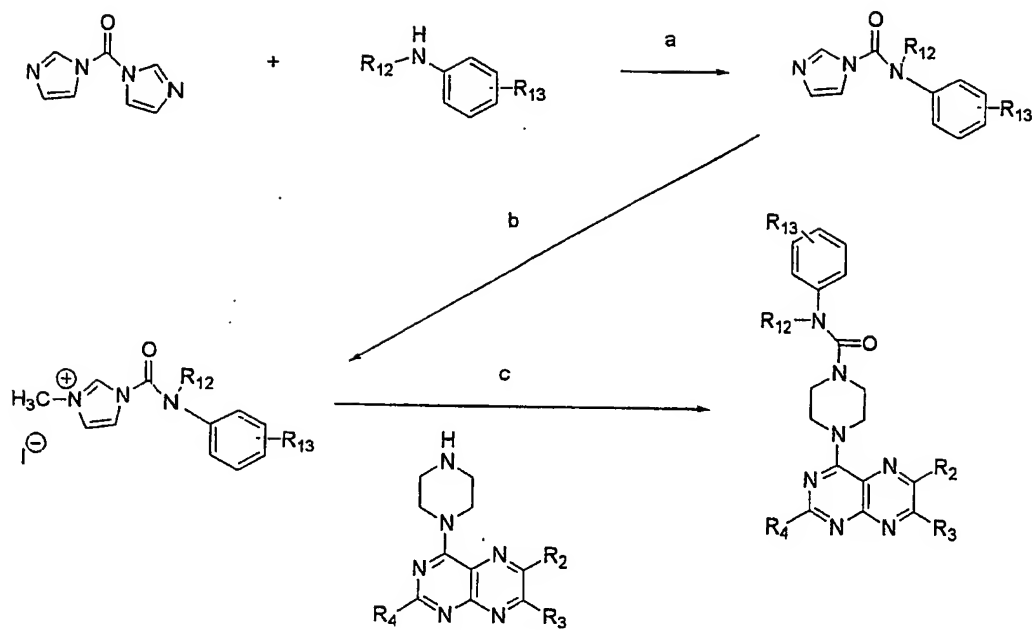
8 / 9

Figure 8



9 / 9

Figure 9



INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP2004/011836

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K31/519 A61K31/551 C07D475/08 A61P37/00 A61P37/06
A61P9/00 A61P25/00 A61P37/08 A61P29/00 A61P31/12
A61P35/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data, PAJ, BIOSIS, CANCERLIT, SCISEARCH, EMBASE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 00/39129 A (WAER MARK JOSEPH ALBERT ; HERDEWIJN PIET ANDRE MAURITS M (BE); LEUVEN) 6 July 2000 (2000-07-06) page 1, line 4 - page 2, line 2 page 7, lines 26-34 page 17, line 30 - page 18, line 13 page 19, line 8 - page 20, line 4 compounds 41-44 claims 1,2,6,15-17	1-7,9, 11-13, 17,18
X	WO 01/21619 A (PFLEIDERER WOLFGANG ; KOTSONIS PETER (DE); SCHMIDT HARALD (DE); FROEHL) 29 March 2001 (2001-03-29) page 1, lines 10-16 page 5, line 1 - page 7, line 10 page 15, lines 11-35 compounds 35-38 claims 1,7,8	1-7,9, 11,17,18

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

18 January 2005

Date of mailing of the international search report

31/01/2005

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Cielen, E

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP2004/011836

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>EP 0 574 906 A (NISSHIN FLOUR MILLING CO) 22 December 1993 (1993-12-22) cited in the application page 4, lines 1-32 page 5, line 47 - page 7, line 57 page 24, lines 51-54 compounds 15,57 claims 1,3,6</p>	<p>1-3,5,7, 11,17,18</p>
X	<p>FROELICH, LOTHAR G. ET AL: "Inhibition of Neuronal Nitric Oxide Synthase by 4-Amino Pteridine Derivatives: Structure-Activity Relationship of Antagonists of (6R)-5,6,7,8-Tetrahydrobiopterin Cofactor" JOURNAL OF MEDICINAL CHEMISTRY, 42(20), 4108-4121 CODEN: JMCMAR; ISSN: 0022-2623, 1999, XP002313877 abstract page 4108, column 2, paragraph 1 tables 9,17; compounds 17FA,17FF page 4119, column 1, paragraph 2</p>	<p>1-5,7,9, 17,18</p>
X	<p>MERZ KARL-HEINZ ET AL: "Synthesis of 7-benzylamino-6-chloro-2-piperazino-4-pyrr olidinopterid ine and novel derivatives free of positional isomers. Potent inhibitors of cAMP-specific phosphodiesterase and of malignant tumor cell growth" JOURNAL OF MEDICINAL CHEMISTRY, vol. 41, no. 24, 19 November 1998 (1998-11-19), pages 4733-4743, XP002313878 ISSN: 0022-2623 abstract table 2; compounds 7G,7I page 4738, column 2, paragraph 4 - page 4739, column 1, paragraph 2 page 4739, column 2, paragraph 4 page 4742, column 1, paragraph 6 page 4742, column 2, paragraph 2</p>	<p>1-3,7,11</p>
X	<p>WEINSTOCK J ET AL: "PTERIDINES. XII. STRUCTURE-ACTIVITY RELATIONSHIPS OF SOME PTERIDINE DIURETICS" JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY. WASHINGTON, US, vol. 11, no. 3, 1968, pages 573-579, XP002908101 ISSN: 0022-2623 abstract page 574, column 1, paragraph 1-3 table VII</p>	<p>1-5,7,9, 11</p>

-/--

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP2004/011836

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2 940 972 A (JOSEF ROCH) 14 June 1960 (1960-06-14) cited in the application column 3, lines 16-20 column 10, lines 28-34 compounds 24-26 claim 1	1-4,7,9, 11
A	Y. LANDRY, J.-P. GIES: "Pharmacologie. Des cibles vers l'indication thérapeutique. Cours et exercices." 2003, DUNOD, PARIS, XP002313503 page 177, paragraphs 2,3	
A	EP 0 185 259 A (THOMAE GMBH DR K) 25 June 1986 (1986-06-25) cited in the application	1,3,5-7
X	page 1, line 11 - page 2, line 24 page 10, lines 1-9 page 13, line 20 - page 15, line 4 examples 1-13 page 22, line 18 - page 23, line 22 examples 14-35 claims 1,7,8	11,17,18
A	EP 0 362 645 A (THOMAE GMBH DR K) 11 April 1990 (1990-04-11) cited in the application page 2, lines 11-43 page 6, lines 21-30 claims 1-4	1,11,17
A	WO 02/32507 A (BONNERT ROGER ; WALTERS IAIN (GB); ASTRAZENECA AB (SE)) 25 April 2002 (2002-04-25) cited in the application page 1, line 31 - page 4, line 20 page 7, line 7 - page 8, line 29	1,11,17
A	NICOLAUS B J R: "Symbiotic Approach to Drug Design" DECISION MAKING IN DRUG RESEARCH, XX, XX, 1983, pages 173-186, XP002197412 the whole document	
T	EP 1 479 682 A (4 AZA BIOSCIENCE NV) 24 November 2004 (2004-11-24) page 2, paragraph 1-3 page 24, paragraphs 67,70,71 page 25, paragraphs 73,74,78,79 page 26, paragraph 82-84 page 27, paragraph 87-91	1,11-18

INTERNATIONAL SEARCH REPORT

International application No.
PCT/EP2004/011836

Box II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claims 17-18 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP2004/011836

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 0039129	A	06-07-2000	AT 277929 T AU 770551 B2 AU 3042900 A CA 2356380 A1 DE 69920757 D1 WO 0039129 A1 EP 1144412 A1 JP 2002533464 T US 2004077859 A1	15-10-2004 26-02-2004 31-07-2000 06-07-2000 04-11-2004 06-07-2000 17-10-2001 08-10-2002 22-04-2004
WO 0121619	A	29-03-2001	DE 19944767 A1 AU 7517400 A WO 0121619 A1 EP 1216246 A1 JP 2004522690 T	29-03-2001 24-04-2001 29-03-2001 26-06-2002 29-07-2004
EP 0574906	A	22-12-1993	HU 64333 A2 CA 2098562 A1 DE 69330626 D1 DE 69330626 T2 EP 0574906 A2 JP 3247769 B2 JP 6179673 A US 5380724 A	28-12-1993 18-12-1993 27-09-2001 04-07-2002 22-12-1993 21-01-2002 28-06-1994 10-01-1995
US 2940972	A	14-06-1960	NONE	
EP 0185259	A	25-06-1986	DE 3445298 A1 AU 576924 B2 AU 5123285 A CA 1252783 A1 DK 572685 A ,B, EP 0185259 A2 ES 8707238 A1 FI 854862 A ,B, GR 852996 A1 IL 77294 A JP 61140585 A NO 854965 A ,B, NZ 214522 A PH 24451 A PT 81650 A ,B ZA 8509462 A	12-06-1986 08-09-1988 19-06-1986 18-04-1989 13-06-1986 25-06-1986 01-10-1987 13-06-1986 16-04-1986 28-02-1989 27-06-1986 13-06-1986 28-07-1988 25-06-1990 01-01-1986 29-07-1987
EP 0362645	A	11-04-1990	DE 3833393 A1 AT 106728 T AU 620645 B2 AU 4174989 A DD 299062 A5 DE 58907824 D1 DK 480489 A EP 0362645 A2 ES 2056174 T3 FI 894615 A ,B, HU 208486 B HU 52504 A2 IL 91814 A JP 2256676 A	05-04-1990 15-06-1994 20-02-1992 05-04-1990 26-03-1992 14-07-1994 02-04-1990 11-04-1990 01-10-1994 02-04-1990 29-11-1993 28-07-1990 27-02-1994 17-10-1990

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/EP2004/011836

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0362645	A	MX 9203034 A1	01-07-1992
		NO 893879 A ,B,	02-04-1990
		NZ 230833 A	28-05-1991
		PT 91836 A ,B	30-04-1990
		SU 1720491 A3	15-03-1992
		ZA 8907414 A	26-06-1991
WO 0232507	A	25-04-2002	
		AU 9615101 A	29-04-2002
		EP 1328319 A1	23-07-2003
		JP 2004511532 T	15-04-2004
		WO 0232507 A1	25-04-2002
EP 1479682	A	24-11-2004	
		US 2003236255 A1	25-12-2003
		EP 1479682 A1	24-11-2004
		WO 2004104005 A2	02-12-2004